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(54) Title: G-PROTEIN FUSION RECEPTORS AND CHIMERIC GABAB RECEPTORS

(57) Abstract

The present invention features G-protein fusion receptors and chimeric GABA_B receptors (GABA_BRs), nucleic acid encoding such receptors, and the use of such receptors and nucleic acid. G-protein fusion receptors comprise at least one domain from a CaR, a mGluR, and/or a GABA_B receptor fused directly or through a linker to a guanine nucleotide-binding protein (G-protein). Chimeric GABA_BRs comprise at least one of a GABA_BR extracellular domain, a GABA_BR transmembrane domain, or a GABA_BR intracellular domain and one or more domains from a mGluR subtype 8 (mGluR8) and/or a CaR.

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DESCRIPTION

G-PROTEIN FUSION RECEPTORS AND CHIMERIC GABA, RECEPTORS

RELATED APPLICATIONS 5

The present application claims priority to Garrett et al. U.S. Serial No. 60/080,671, filed April 3, 1998, which is hereby incorporated by reference herein in its entirety including the drawings.

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FIELD OF THE INVENTION

The present invention relates to a G-protein fusion receptors, chimeric GABA, (y-aminobutyric acid) receptors, nucleic acid encoding such receptors, and uses of such receptors and nucleic acid encoding such receptors.

BACKGROUND

The references cited herein are not admitted to be prior art to the claimed invention.

Chimeric receptors made up of peptide segments from different receptors have different uses such as being used to assess the functions of different sequence regions and to assess the activity of different compounds at a particular receptor. Examples of using chimeric receptors to assess the activity of 25 different compounds are provided by Dull et al., U.S. Patent No. 4,859,609, Dull et al., U.S. Patent No. 5,030,576, and Fuller et al., International Application No. PCT/US96/12336, International Publication No. WO 97/05252.

Dull et al. U.S. Patent No. 4,859,609, and Dull et al. U.S. Patent No. 5,030,576, indicate the production and use of chimeric receptors comprising a ligand binding domain of a predetermined receptor and a heterologous reporter polypeptide. The Dull et al. patents provide as examples of chimerics: (1) a chimeric receptor made up of the insulin receptor extracellular α chain, and the EGF receptor transmembrane and cytoplasmic domains without any HIR B-chain sequence; and (2) a hybrid receptor made

up of the v-erB oncogene product intracellular domain fused to the EGF receptor extracellular and transmembrane domains.

Fuller et al. International Publication No. WO 97/05252 feature chimeric receptors made up of metabotropic glutamate receptor (mGluR) domains and calcium receptor (CaR) domains. The chimeric receptors allow the coupling of functional aspects of a mGluR with a CaR.

An example of the use of chimeric receptors to assess the functions of different sequence regions receptors are found in studies identifying regions of different guanine nucleotide-binding protein coupled receptors important for guanine nucleotide-binding protein coupling. (See, Kobilka et al., Science 240:1310-1316, 1988; Wess et al., FEBS Lett. 258:133-136, 1989; Cotecchia et al., Proc. Natl. Acad. Sci. USA 87:2896-2900, 1990; Lechleiter et al., EMBO J. 9:4381-4390, 1990; Wess et al., Mol. Pharmacol. 38:517-523, 1990; and Pin et al., EMBO J. 13:342-348, 1994.)

SUMMARY OF THE INVENTION

- The present invention features G-protein fusion receptors and chimeric GABA_B receptors (GABA_BRs), nucleic acid encoding such receptors, and the use of such receptors and nucleic acid. G-protein fusion receptors comprise at least one domain from a CaR, a mGluR, and/or a GABA_B receptor fused directly or through a linker to a guanine nucleotide-binding protein (G-protein). Chimeric GABA_BRs comprise at least one of a GABA_BR extracellular domain, a GABA_BR transmembrane domain, or a GABA_BR intracellular domain and one or more domains from a mGluR subtype 8 (mGluR8) and/or a CaR.
- G-proteins are peripheral membrane proteins made up of an α subunit, a β subunit, and a γ subunit. G-proteins interconvert between a GDP bound and a GTP bound form. Different types of G-proteins can affect different enzymes, such as adenylate cyclase and phospholipase-C.
- 35 Thus, a first aspect of the present invention describes a Gprotein fusion receptor comprising:

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an extracellular domain comprising an extracellular domain amino acid sequence substantially similar to either an extracellular CaR amino acid sequence, an extracellular mGluR amino acid sequence, or an extracellular GABAB receptor amino acid sequence;

a transmembrane domain joined to the carboxy terminus of said extracellular domain, said transmembrane domain comprising a transmembrane domain amino acid sequence substantially similar to either a transmembrane CaR amino acid sequence, a transmembrane mGluR amino acid sequence, or a transmembrane GABA_B receptor amino acid sequence;

an intracellular domain joined to the carboxy terminus of said transmembrane domain comprising all or a portion of an intracellular amino acid sequence substantially similar to either an intracellular CaR amino acid sequence, an intracellular mGluR amino acid sequence, or an intracellular GABA $_{B}$ receptor amino acid sequence, provided that said portion is at least about 10 amino acids;

an optionally present linker joined to the carboxy terminus of said intracellular domain; and

a G-protein joined either to said intracellular domain or to said optionally present linker, provided that said G-protein is joined to said optionally present linker when said optionally present linker is present.

"Substantially similar" refers to at least 40% sequence similarity between respective polypeptide regions making up a domain. In preferred embodiments, substantially similar refers to at least 50%, at least 75%, at least 90%, at least 95% sequence similarity, or 100% (the same sequence), between polypeptide domains. The degree to which two polypeptide domains are substantially similar is determined by comparing the amino acid sequences located in corresponding domains. Sequence similarity is preferably determined using BLASTN (Altschul et al., J. Mol. Biol. 215:403-410, 1990).

The different receptor components of the G-protein receptor can come from the same receptor protein or from a chimeric receptor made up of different receptor domains. By swapping different domains compounds able to effect different domains of a

particular receptor can be identified and the activity of different compounds at different domains can be measured.

In different embodiments the CaR region(s) present in the G-protein fusion are substantially similar to, comprise, or consist of portion(s) of the human CaR; mGluR region(s) present in the G-protein fusion are substantially similar to, comprise, or consist of portion(s) of a human mGluR; and GABA_BR region(s) present in the G-protein fusion are substantially similar to, comprise, or consist of portion(s) of a human GABA_BR receptor.

In preferred embodiments concerning GABA_ER regions that are present: the GABA_ER extracellular domain is substantially similar to a GABA_ER extracellular domain provided in SEQ. ID. NOs. 2-4; the GABA_ER transmembrane domain is substantially similar to the GABA_ER transmembrane domain provided in SEQ. ID. NOs. 7-9; and the GABA_ER intracellular domain is substantially similar to a GABA_ER intracellular domain provided in SEQ. ID. NOs. 12-14.

In preferred embodiments concerning CaR regions that are present: the CaR extracellular domain is substantially similar to the CaR extracellular provided in SEQ. ID. NO. 1; the CaR transmembrane domain is substantially similar to the CaR transmembrane domain provided in SEQ. ID. NO. 6; and the CaR intracellular domain is substantially similar to the CaR intracellular domain such as that provided in SEQ. ID. NO. 11.

Various different mGluR subtypes present in different organisms, including humans, are described in different patent publications as follows: mGluR₁ - WO 94/29449, EP 569 240 Al, WO 92/10583 and U.S. Patent No. 5,385,831; mGluR₂ - WO 94/29449, WO 96/06167, and EP 711 832 A2; mGluR₃ - WO 94/29449, and WO 95/22609; mGluR₄ - WO 95/08627, WO 95/22609, and WO 96/29404; 30 mGluR₅ - WO 94/29449; mGluR₆ - WO 95/08627; mGluR₇ - U.S. Patent No. 5,831,047, WO 95/08627 and WO 96/29404; and mGluR₆ - WO 97/48724 and EP 616 498 A2. (Each of these references are hereby incorporated by reference herein.)

In preferred embodiments concerning mGluR regions that are present: the mGluR extracellular domain is substantially similar to either human mGluR 1, human mGluR 2, human mGluR 3, human mGluR 4, human mGluR 5, human mGluR 6, human mGluR 7, or human

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mGluR 8; the mGluR transmembrane domain is substantially similar to either human mGluR 1, human mGluR 2, human mGluR 3, human mGluR 4, human mGluR 5, human mGluR 6, human mGluR 7, or human mGluR 8; and the mGluR intracellular domain is substantially similar to either human mGluR 1, human mGluR 2, human mGluR 3, human mGluR 4, human mGluR 5, human mGluR 6, human mGluR 7, or human mGluR 8.

Another aspect of the present invention describes a nucleic acid comprising a nucleotide sequence encoding for a G-protein fusion receptor.

Another aspect of the present invention describes a recombinant cell comprising an expression vector encoding for a G-protein fusion receptor, and a cell where the G-protein fusion receptor is expressed. Preferably, the G-protein fusion receptor is functional in the cell.

Another aspect of the present invention describes a recombinant cell produced by combining (a) a cell where a G-protein fusion receptor is expressed, and (b) a vector comprising nucleic acid encoding a G-protein fusion receptor and elements for introducing heterologous nucleic acid into the cell. Preferably, the G-protein fusion receptor is functional in the cell.

Another aspect of the present invention describes a process for the production of a G-protein fusion receptor. The process is performed by growing host cells comprising a G-protein fusion receptor.

Another aspect of the present invention describes a method of measuring the ability of a compound to affect G-protein fusion receptor activity.

Another aspect of the present invention describes a chimeric GABA_BR comprising an extracellular domain, a transmembrane domain and an intracellular domain, wherein at least one domain is from a GABA_BR and at least one domain is from CaR or mGluR8. The extracellular domain comprises an amino acid sequence substantially similar to a CaR extracellular domain (SEQ. ID. NO.

- 1), a GABA_BRla extracellular domain (SEQ. ID. NO. 2), a GABA_BRlb extracellular domain (SEQ. ID. NO. 3), a GABA_BR2 extracellular domain (SEQ. ID. NO. 4), or a mGluRB extracellular domain (SEQ. ID. NO. 5).
- The transmembrane domain comprises an amino acid sequence substantially similar to a CaR transmembrane domain (SEQ. ID. NO. 6), a GABA_BRla transmembrane domain (SEQ. ID. NO. 7), a GABA_BRlb transmembrane domain (SEQ. ID. NO. 8), a GABA_BR2 transmembrane domain (SEQ. ID. NO. 9), or a mGluR8 transmembrane domain (SEQ. ID. NO. 10).

The intracellular domain comprises an amino acid sequence substantially similar to a CaR intracellular domain (SEQ. ID. NO. 11), a GABA_BR1a intracellular domain (SEQ. ID. NO. 12), a GABA_BR1b intracellular domain (SEQ. ID. NO. 13), a GABA_BR2 intracellular domain (SEQ. ID. NO. 13), or a mGluR8 intracellular domain (SEQ. ID. NO. 15).

Preferred chimeric $GABA_gRs$ contain at least one mGluR8 intracellular, transmembrane or extracellular domain, or at least one CaR intracellular, transmembrane or extracellular domain.

More preferably, the chimeric $GABA_{\tt p}R$ contains at least one CaR domain.

In preferred embodiments concerning mGluR8 regions that are present: the mGluR8 extracellular domain is substantially similar to the mGluR8 extracellular domain provided in SEQ. ID. NO. 5; the mGluR8 transmembrane domain is substantially similar to the mGluR8 transmembrane domain provided in SEQ. ID. NO. 10; and the mGluR8 intracellular domain is substantially similar to the mGluR8 receptor intracellular provided in SEQ. ID. NO. 15.

Preferably, the domains are functionally coupled such that a signal from the binding of an extracellular ligand is transduced to the intracellular domain when the chimeric receptor is present in a suitable host cell. A suitable host cell contains the elements for functional signal transduction for receptors coupled to a G-protein.

Another aspect of the present invention describes a nucleic acid comprising a nucleotide sequence encoding for a chimeric $GABA_BR$.

Another aspect of the present invention describes a recombinant cell comprising an expression vector encoding for a chimeric $GABA_BR$, and a cell where the chimeric $GABA_BR$ is expressed. Preferably, the chimeric $GABA_BR$ is functional in the cell.

Another aspect of the present invention describes a recombinant cell produced by combining (a) a cell where a chimeric $GABA_pR$ is expressed, and (b) a vector comprising nucleic acid encoding the chimeric $GABA_pR$ and elements for introducing heterologous nucleic acid into the cell. Preferably, the chimeric $GABA_pR$ is functional in the cell.

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Another aspect of the present invention describes a process for the production of a chimeric receptor. The process is performed by growing host cells comprising a chimeric GABA_BR.

Another aspect of the present invention describes a method of measuring the ability of a compound to affect $GABA_BR$ or mGluR activity. The method is performed by measuring the ability of a compound to affect chimeric $GABA_BR$ or mGluR activity.

Another aspect of the present invention describes a fusion receptor polypeptide comprising a receptor and a G-protein α subunit, wherein said G-protein α subunit is fused to the intracellular domain of said receptor, provided that the receptor is not an adrenoreceptor.

Various examples are described herein. These examples are not intended in any way to limit the claimed invention.

Other features and advantages of the invention will be apparent from the following drawings, the description of the invention, the examples, and the claims.

BRIEF DESCRIPTION OF DRAWINGS

Figures 1a-1d illustrate the amino acid sequences of a human CaR extracellular domain (SEQ. ID. NO. 1), a human GABA $_{\rm B}$ R1a extracellular domain (SEQ. ID. NO. 2), a human GABA $_{\rm B}$ R1b extracellular domain (SEQ. ID. NO. 3), a human GABA $_{\rm B}$ R2 extracellular domain (SEQ. ID. NO. 4), and a human mGluR8 extracellular domain (SEQ. ID. NO. 5).

Figures 2a-2b illustrate the amino acid sequences of a human CaR transmembrane domain (SEQ. ID. NO. 6), a human GABA_BRla transmembrane domain (SEQ. ID. NO. 7), a human GABA_BRlb transmembrane domain (SEQ. ID. NO. 8), a human GABA_BR2 transmembrane domain (SEQ. ID. NO. 9), and a human mGluR8 transmembrane domain (SEQ. ID. NO. 10).

Figures 3a-3b illustrate the amino acid sequences of a human CaR intracellular domain (SEQ. ID. NO. 11), a human GABABRIA intracellular domain (SEQ. ID. NO. 12), a human GABABRIb intracellular domain (SEQ. ID. NO. 13), a human GABABR2 intracellular domain (SEQ. ID. NO. 14), and a human mGluR8 intracellular domain (SEQ. ID. NO. 15).

20 Figures 4a-4b illustrate the amino acid sequence of $G_{\alpha_{15}}$ (SEQ. ID. No. 16) and $G_{\alpha_{16}}$ (SEQ. ID. No. 17).

Figures 5a-5r illustrate the cDNA sequences encoding for human CaR (SEQ. ID. NO. 18), human GABA $_{\rm e}$ R1a (SEQ. ID. NO. 19), human GAEA $_{\rm e}$ R1b (SEQ. ID. NO. 20), and human GABA $_{\rm e}$ R2 (SEQ. ID. NO.

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Figures 6a-6h illustrate the cDNA sequence for rat GABA,Rla (SEO. ID. NO. 22) and rat GABA,Rlb (SEQ. ID. NO. 23).

Figures 7a-7c illustrate the amino sequence for rat GABA_R1a (SEQ. ID. NO. 24) and rat GABA_R1b (SEQ. ID. NO. 25).

Figure 8 illustrates the ability of a chimeric CaR/GABA_BR2 (CaR extracellular and transmembrane domains, and intracellular GABA_BR2 domain) to transduce a signal. Signal production was measured by detecting an increase in the calcium-activated chloride current. The line in the middle of the increase signifies a wash step.

Figures 9a-9p illustrate the cDNA sequence for human mGluR2

(SEQ. ID. NO. 26), chimeric hCAR/hmGluR2 (SEQ. ID. NO. 30), chimeric hmGluR2/hCaR (SEQ. ID. NO. 34), and chimeric hmGluR8/hCaR (SEQ. ID. NO. 38).

Figures 10a-10f illustrate the amino acid sequence for human 5 mGluR2 (SEQ. ID. NO. 27), chimeric hCAR/hmGluR2 (SEQ. ID. NO. 31), chimeric hmGluR2/hCaR (SEQ. ID. NO. 35), chimeric hmGluR8/hCaR (SEQ. ID. NO. 39).

Figures lla-11v illustrate the cDNA sequence for the phCaR/hmGluR2*Gqi5 fusion construct (SEQ. ID. NO. 32), pmGluR2//CaR*G α_q i5 fusion construct (SEQ. ID. NO. 36), pmGluR2//CaR*G α_q i5+3Ala linker fusion construct (SEQ. ID. NO. 10 46), and the mGluR8//CaR*G α_q i5 fusion construct (SEQ. ID. NO. 40).

Figures 12a-12h illustrate the amino acid sequence for the phCaR/hmGluR2*Gqi5 fusion construct (SEQ. ID. NO. 33), pmGluR2//CaR*G α_q i5 fusion construct (SEQ. ID. NO. 37), 15 pmGluR2//CaR*G α_q i5+3Ala linker fusion construct (SEQ. ID. NO. 47), and the mGluR8//CaR*G α_q i5 fusion construct (SEQ. ID. NO. 41).

Figures 13a-13m illustrate the cDNA sequence for the GABA-R2*Gqo5 fusion construct (SEQ. ID. NO. 42) and the GABA-BRla*Gqo5 20 fusion construct (SEQ. ID. NO. 44).

Figures 14a-14e illustrates the amino acid sequence for the GABA-BR2+Gqo5 fusion construct (SEQ. ID. NO. 43) and the GABA-BR1a*Gq05 fusion construct (SEQ. ID. NO. 45).

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Figure 15 illustrates the ability of different G-protein fusions to transduce signal resulting from ligand binding. mGluR2//CaR'Gqi5 is shown by SEQ. ID. NO. 37, CaR/mGluR2'Gqi5 is shown by SEQ. ID. NO. 33, mGluR8//CaR*Gqi5 is shown by SEQ. ID. NO. 41.

DETAILED DESCRIPTION OF THE INVENTION

The CaR, mGluR, and the $GABA_{\theta}R$ are structurally similar in that they are each a single subunit membrane protein possessing an extracellular domain, a transmembrane domain comprising seven putative membrane spanning helices connected by three intracellular and three extracellular loops, and an intracellular carboxy-terminal domain. Signal transduction is activated by the extracellular binding of an agonist. The signal is transduced to the intracellular components of the receptor causing an intracellular effect.

Signal transduction from agonist binding to an extracellular region can be modulated by compounds acting at a downstream transmembrane domain or the intracellular domain. Downstream effects include antagonist actions of compounds and allosteric actions of compounds.

The transmembrane domain provides different types of target sites for compounds modulating receptor activity in different environments. As noted above, the transmembrane domain contains extracellular, transmembrane, and intracellular components.

Compounds modulating GABA_ER, CaR, or mGluR activity can be obtained, for example, by screening a group or library of compounds to identify those compounds having the desired activity and then synthesizing such compound. Thus, included in the present invention is a method of making a GABA_BR, CaR, or mGluR active compound by first screening for a compound having desired properties and then chemically synthesizing that compound.

Metabotropic Glutamate Receptors (mGluRs)

mGluRs are G protein-coupled receptors capable of activating a variety of intracellular secondary messenger systems following the binding of glutamate (Schoepp et al., Trends Pharmacol. Sci. 11:508, 1990; Schoepp and Conn, Trends Pharmacol. Sci. 14:13, 1993, hereby incorporated by reference herein).

Activation of different mGluR subtypes in situ elicits one or more of the following responses: activation of phospholipase C, increases in phosphoinositide (PI) hydrolysis, intracellular calcium release, activation of phospholipase D, activation or inhibition of adenylyl cyclase, increases and decreases in the formation of cyclic adenosine monophosphate (cAMP), activation of guanylyl cyclase, increases in the formation of cyclic guanosine monophosphate (cGMP), activation of phospholipase A₁, increases in

arachidonic acid release, and increases or decreases in the activity of voltage- and ligand-gated ion channels (Schoepp and Conn, Trends Pharmacol. Sci. 14:13, 1993; Schoepp, Neurochem. Int. 24:439, 1994; Pin and Duvoisin, Neuropharmacology 34:1, 1995, hereby incorporated by reference herein).

Eight distinct mGluR subtypes have been isolated. (Nakanishi, Neuron 13:1031, 1994; Pin and Duvoisin, Neuropharmacology 34:1, 1995; Knopfel et al., J. Med. Chem. 38:1417; Eur. J. Neuroscience 7:622-629, 1995, each of these references is hereby incorporated by reference herein.) The different mGluRs possess a large aminoterminal extracellular domain (ECD) followed by seven putative transmembrane domain (7TMD) comprising seven putative membrane spanning helices connected by three intracellular and three extracellular loops, and an intracellular carboxy-terminal domain of variable length (cytoplasmic tail). 15

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Human mGluR8 is described by Stormann et al., International Application Number PCT/US97/09025, International Publication Number WO 97/48724, and mouse mGluR8 is described by Duvoisin et al., J. Neurosci. 15:3075-3083, 1995, (both of these references are hereby incorporated by reference herein). mGluR8 couples to G_{i} . Agonists of mGluR8 include L-glutamate and L-2-amino-4phosphonobutyrate.

mGluR8 activity can be measured using standard techniques. For example, G_i negatively couples to adenylate cyclase to inhibit intracellular cAMP accumulation in a pertussis toxin-sensitive fashion. Thus, mGluR8 activity can be measured, for example, by measuring inhibition of forskolin-stimulated cAMP production as described by Duvoisin et al., J. Neurosci. 15:3075-3083, 1995.

mGluRs have been implicated in a variety of neurological pathologies. Examples of such pathologies include stroke, head trauma, spinal cord injury, epilepsy, ischemia, hypoglycemia, 30 anoxia, and neurodegenerative diseases such as Alzheimer's disease (Schoepp and Conn, Trends Pharmacol. Sci. 14:13, 1993; Cunningham et al., Life Sci. 54: 135, 1994; Pin et al., Neuropharmacology 34:1, 1995; Knopfel et al., J. Med. Chem.

38:1417, 1995, each of which is hereby incorporated by reference herein).

Calcium Receptor

The CaR responds to changes of extracellular calcium concentration and also responds to other divalent and trivalent cations. The CaR is a G-protein coupled receptor containing an extracellular Ca2+ binding domain. Activation of the CaR, descriptions of CaRs isolated from different sources, and examples of CaR active compound are provided in Nemeth NIPS 10:1-10 5, 1995, Brown et al. U.S. Patent No. 5,688,938, Van Wagenen et al., International Application Number PCT/US97/05558 International Publication Number WO 97/37967, Brown E.M. et al., Nature 366:575, 1993, Riccardi D., et al., Proc. Nat'l. Acad. 15 Sci. USA 92:131-135, 1995, and Garrett J.E., et al., J. Biol. Chem. 31:12919-12925, 1995. (Each of these references are hereby incorporated by reference herein.) Brown et al. U.S. Patent No. 5,688,938 and Van Wagenen et al., International Application Number PCT/US97/05558 International Publication Number WO 97/37967, describe different types of compounds active at the CaR 20 including compounds which appear to be allosteric modulators and CaR antagonists.

The CaR can be targeted to achieve therapeutic effects.

Examples of target diseases are provided in Brown et al. U.S.

Patent No. 5,688,938, and Van Wagenen et al., International Application Number PCT/US97/05558 International Publication Number WO 97/37967, and include hyperparathyroidism and osteoporosis.

y-Aminobutyric acid Receptors (GABA_BRs)

GABA_ERs are G-protein coupled metabotropic receptors. GABA_ERs modulate synaptic transmission by inhibiting presynaptic transmitter release and by increasing K conductance responsible for long-lasting inhibitory postsynaptic potentials. (See,

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Kaupmann et al., Nature 386:239-246, 1997, hereby incorporated by reference herein.)

GABA.Rs are found in the mammalian brain, in locations outside of the brain, and in lower species. Outside of the brain, GABA,Rs have been identified on axon terminals and ganglion cell bodies of the autonomic nervous system, on fallopian tube and uterine intestinal smooth muscle cells, in the kidney cortex, urinary bladder muscle and on testicular interstitial cells. (See, Bowery, Annu. Rev. Pharmacol. Toxicol. 33:109-147, 1993, hereby incorporated by reference herein.)

Different GABA, Rs subtypes exist. Kaupmann et al., Nature 386:239-246, 1997, indicate that they cloned GABA,Rs. Nucleic acid encoding two $GABA_{e}R$ proteins were indicated to be cloned from rat brain: GABA,Rla and GABA,Rlb. GABA,Rla differs from GABA,Rlb in that the N-terminal 147 residues are replaced by 18 amino acids. $GABA_BR1a$ and $GABA_BR1b$ appear to be splice variants. The cloned GABA,Rs were indicated to negatively couple adenylyl cyclases and show sequence similarity to the metabotropic receptors for L-glutamate (mGluR). Northern blot analysis 20 indicated that $GABA_gR1a$ and $GABA_gR1b$ is present in brain and testis, but not in kidney, skeletal muscle, liver, lung, spleen, or heart.

Kaupmann et al., International Application Number PCT/EP97/01370, International Publication Number WO 97/46675, 25 indicate that they have obtained rat GABABR clones, GABABRIa and GABABRID; and humans GABABR clones, GABABRIa/b (representing a partial receptor clone) and GABA,R1b (representing a full-length receptor clone). Amino acid sequence information, and encoding cDNA sequence information, is provided for the different $\textsc{GABA}_{\mathtt{R}}R$ 30 clones.

Another GABA,R subtype is GABA,R2. Northern blot analysis reveals than an approximately 6.3 Kb human GABA,R2 transcript is abundantly expressed in the human brain. Expression is not detected in the heart, placenta, lung, liver, skeletal muscle, kidney and pancreas under conditions where GABA_BR2 transcript was identified in the human brain. Within the human brain $GAEA_{\mu}R2$ is broadly expressed at variable levels.

GABA_BR functions as a heterodimer of the subunits GABA_BRl or GABA_BR2. (Jones et al. Nature 396:674-679, 1998, hereby incorporated by reference herein.)

GABA_BRs have been targeted to achieve therapeutic effects.

Kerr and Ong, DDT 1:371-380, 1996, describe different compounds indicated to be GABA_BR agonists and GABA_BR antagonists. Kerr and Ong also review therapeutic implications of affecting GABA_BR activity including, spasticity and motor control, analgesia, epilepsy, cognitive effects, psychiatric disorders, alcohol dependence and withdrawal, feeding behavior, cardiovascular and respiratory functions, and peripheral functions.

Bittiger et al., Tips 4:391-394, 1993, review therapeutic applications of GABA_BR antagonists. Potential therapeutic applications noted by Bittiger et al. include cognitive processes, epilepsy, and depression.

G-Protein Fusion Receptors

Examples of some different types of G-protein fusion receptors, and advantages of some receptors, are provided below.

Using the present application as guide additional G-protein receptors fusion can be constructed.

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G-protein fusion receptors contain an intracellular domain of a receptor fused to a G-protein α subunit (G_{α}) . G_{α} fusions to adrenoreceptors have been reported by Bertin et al., Receptors and Channels 5:41-51, 1997; Wise and Milligan, Journal of Biological Chemistry 39:24673-24678, 1997; and Bertin et al., Proc. Natl. Acad. Sci. USA 91:8827-8831, 1994 (each of which are hereby incorporated by reference herein). These studies were indicated to produce a functional chimeric by fusing the α_{2k} -adrenoreceptor to the $G_{3\alpha}$, or the β_{1} -adrenoreceptor to the $G_{5\alpha}$.

The G-protein fusion receptors described by the present invention include a G-protein fused to an intracellular domain, where the intracellular domain when present in a wild type

receptor does not interact with that type of G-protein. Thus, the present invention also describes swapping of signals by fusing an intracellular domain to a G_{α} normally not coupled to that intracellular domain. The use of such fusion proteins, while applicable to chimeric GABA_BRs, is not limited to chimeric GABA_BRs. Indeed, such technology can be applied to receptors containing an extracellular domain, transmembrane domain and intracellular domain of a wild type receptor.

Preferred G-proteins fusion receptors contain an intracellular domain fused to a promiscuous G_{α} that couples to phospholipase C resulting in the mobilization of intracellular calcium. Increases in intracellular calcium can be conveniently measured through the use of dyes. Such techniques are well known in the art and are described, for example by Brown et al. U.S. Patent No. 5,688,938.

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In an embodiment G-proteins fusions can also be used to decrease receptor desensitization.

Examples of promiscuous G_{α} 's coupling to phospholipase C include naturally occurring G-proteins such as $G_{\alpha_{15}}$ and $G_{\alpha_{16}}$, and chimeric G-protein such as Gqo5 and Gqi5. Gqo5 and Gqi5 are made of a Gq portion where the five amino acids at the C-terminal are from either G_{\circ} or G_{\circ} respectively (Conklin et al., Nature 363:274-277, 1993, hereby incorporated by reference herein). The Gq portion of such chimeric receptors provides for phospholipase C coupling while the terminal G_{\circ} or G_{\circ} portion allows the chimeric G-protein to couple to different receptor proteins that are normally involved in inhibitor effects on adenylate cyclase.

In an embodiment of the present invention the employed G-protein is from a human source or is made up of different G-protein components each from a human source.

G-proteins fusions can be created, for example, by fusing directly or indirectly the intracellular domain of a receptor protein to a polypeptide having an amino acid sequence substantially similar to $G_{\Omega 15}$, $G_{\Omega 16}$, G_{Q05} or G_{Q15} . In different embodiments, the receptor is fused directly or indirectly to a G-

protein consisting of the amino acid sequence of $G_{\alpha_{15}},\ G_{\alpha_{16}},\ G_{qo5}$ or Gqi5.

The intracellular domain portion of a receptor protein fused directly or indirectly to a G-protein should be at least about 10 amino acids in length. In different embodiments the portion is at least about 50 amino acids, at least about 100 amino acids, or the full length of an intracellular domain.

The intracellular domain can be directly linked to a Gprotein or can be indirectly linked through an optionally present
linker. Optionally present linkers are preferably about 3 to
about 30 amino acids in length. Preferred linkers are made up of
alanine, glycine, or a combination thereof.

Chimeric Receptors

Examples of some different types of chimeric receptors, and advantages of some receptors, are provided below. Using the present application as guide additional chimeric receptors can be constructed.

20 Chimeric GABA, R Extracellular Domain

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Chimeric GABA_BRs containing a GABA_BR extracellular domain are particularly useful for studying the importance of the GABA_BR extracellular domain and assaying for compounds active at the extracellular domain. Preferably chimeric GABA_BRs containing a GABA_BR extracellular domain also contain a CaR intracellular domain.

A variety of different activities have been generally attributed to GABA $_{B}$ R subtypes. (E.g., Kerr and Ong, DDT 1:371-380, 1996.) Kaupmann et al., Nature 386:239-246, 1997, report that in preliminary experiments involving GABA $_{B}$ Rla they did not detect positive coupling to the adenylyl cyclase or coupling to the phospholipase effector system.

An intracellular CaR domain can be used to couple with Gproteins which activate phospholipase C and mobilize intracellular calcium. Mobilization of intracellular calcium is

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readily detected, for example, by fluorescent indicators of intracellular Ca2.

An additional advantage of using the intracellular CaR domain is that CaR G-protein activation is not rapidly 5 desensitized. Thus, the intracellular CaR domain can be used to produce a stronger intracellular signal than a signal produced from a receptor which is desensitized rapidity.

More preferably, the chimeric $GABA_BR$ contains an intracellular CaR domain, and also contains either a CaR or a 10 $GABA_BR$ transmembrane domain. Advantages of using a CaR transmembrane domain include separating the effects occurring at a $GABA_BR$ extracellular domain from effects occurring at a transmembrane domain; and providing additional intracellular elements, present on transmembrane intracellular loops, useful for coupling to G-protein.

A $GABA_BR$ transmembrane domain is useful for examining whether the transmembrane $GABA_{\scriptscriptstyle B}R$ can be targeted to affect $GABA_{\scriptscriptstyle B}R$ activity, and obtaining compounds active at the $\mbox{GABA}_{\mbox{\tiny B}}\mbox{R}$ transmembrane domain. For example, a transmembrane ${\tt GABA_BR}$ can be used to screen for transmembrane allosteric modulators and antagonists.

Chimeric GABA, R Transmembrane Domain

Chimeric GABA,Rs containing a GABA,R transmembrane are particularly useful for studying the importance of the $\mbox{\scriptsize GABA}_BR$ transmembrane domain and assaying for compounds active at the transmembrane domain. Preferably Chimeric $GABA_BRs$ containing a $GABA_{\epsilon}R$ transmembrane domain contain an extracellular domain which is either mGluR8 or CaR, and an intracellular CaR domain.

More preferably, the chimeric $GABA_BR$ contains an extracellular domain from either mGluR8 or CaR, a GABA $_{\!B}R$ transmembrane, and an intracellular CaR domain. A chimeric $GABA_{g}R$ containing extracellular mGluR8 or CaR domains can readily be stimulated using mGluR8 or CaR ligands.

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Chimeric GABA,R Intracellular Domain

Chimeric GABA_BRs containing a GABA_ER intracellular domain are particularly useful for studying the importance of the GABA_ER intracellular domain and assaying for compounds active at the intracellular domain. Preferably, the chimeric receptors contain an extracellular domain from either mGluR8 or CaR. The extracellular mGluR8 or CaR domains can readily be activated using mGluR8 or CaR ligands.

10 Receptor Domains

Domains of a G-protein fusion receptor, a chimeric receptor, and G_{Ω} , substantially similar to a particular sequence can be readily produced using the disclosure provided herein in conjunction with information well known in the art. Substantially similar sequences can be obtained taking into account sequence information for a particular type of receptor obtained from different sources, different types of amino acids which are to some extent interchangeable, and the ease of experimentation with which functional receptor activity can be assayed.

Substantially similar sequences includes amino acid alterations such as deletions, substitutions, additions, and amino acid modifications. A "deletion" refers to the absence of one or more amino acid residue(s) in the related polypeptide. An "addition" refers to the presence of one or more amino acid residue(s) in the related polypeptide. Additions and deletions to a polypeptide may be at the amino terminus, the carboxy terminus, and/or internal. Amino acid "modification" refers to the alteration of a naturally occurring amino acid to produce a non-naturally occurring amino acid. A "substitution" refers to the replacement of one or more amino acid residue(s) by another amino acid residue(s) in the polypeptide. Derivatives can contain different combinations of alterations including more than one alteration and different types of alterations.

The sequences of polypeptides can be compared from different sources to help identify variable amino acids not essential for receptor activity. For example, Figure 7 provides the rat GABABRIa and GABABRIB amino acid sequences. The rat GABABRIA and GABABRIB amino acid sequences can be compared with the human GABABRIA and GABABRIB sequences to identify conserved and variable amino acids. Derivatives can then be produced where a variable amino acid is changed, and receptor activity can be readily tested.

Similarly, the amino acid sequences for CaR, mGluR8, and G-proteins from different sources are either known in the art or can readily be obtained. Examples of such references are provided above.

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While the effect of an amino acid change varies depending upon factors such as phosphorylation, glycosylation, intra-chain 15 linkages, tertiary structure, and the role of the amino acid in the active site or a possible allosteric site, it is generally preferred that a substituted amino acid is from the same group as the amino acid being replaced. To some extent the following groups contain amino acids which are interchangeable: the basic 20 amino acids lysine, arginine, and histidine; the acidic amino acids aspartic and glutamic acids; the neutral polar amino acids serine, threonine cysteine, glutamine, asparagine and, to a lesser extent, methionine; the nonpolar aliphatic amino acids 25 glycine, alanine, valine, isoleucine, and leucine (however, because of size, glycine and alanine are more closely related and valine, isoleucine and leucine are more closely related); and the aromatic amino acids phenylalanine, tryptophan, and tyrosine. In addition, although classified in different categories, alanine, glycine, and serine seem to be interchangeable to some extent, 30 and cysteine additionally fits into this group, or may be classified with the polar neutral amino acids.

While proline is a nonpolar neutral amino acid, its replacement represents difficulties because of its effects on conformation. Thus, substitutions by or for proline are not

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preferred, except when the same or similar conformational results can be obtained. The conformation conferring properties of proline residues may be obtained if one or more of these is substituted by hydroxyproline (Hyp).

Examples of modified amino acids include the following: altered neutral nonpolar amino acids such as ω -amino acids of the formula $H_2N(CH_2)_nCOOH$ where n is 2-6, sarcosine (Sar), t-butylalanine (t-BuAla), t-butylglycine (t-BuGly), N-methyl isoleucine (N-MeIle), and norleucine (Nleu); altered neutral aromatic amino acids such as phenylglycine; altered polar, but neutral amino acids such as citrulline (Cit) and methionine sulfoxide (MSO); altered neutral and nonpolar amino acids such as cyclohexyl alanine (Cha); altered acidic amino acids such as cysteic acid (Cya); and altered basic amino acids such as ornithine (Orn).

Preferred derivatives have one or more amino acid alteration(s) which do not significantly affect the receptor activity of the related receptor protein. In regions of receptor domains not necessary for receptor activity, amino acids may be deleted, added or substituted with less risk of affecting activity. In regions required for receptor activity, amino acid alterations are less preferred as there is a greater risk of affecting receptor activity.

Derivatives can be produced using standard chemical

techniques and recombinant nucleic acid techniques.

Modifications to a specific polypeptide may be deliberate, as through site-directed mutagenesis and amino acid substitution during solid-phase synthesis, or may be accidental such as through mutations in hosts which produce the polypeptide.

Polypeptides including derivatives can be obtained using standard techniques such as those described by Sambrook et al., Molecular Cloning, Cold Spring Harbor Laboratory Press (1989). For example, Chapter 15 of Sambrook describes procedures for site-directed mutagenesis of cloned DNA.

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Receptor Nucleic Acid

G-protein fusion and chimeric receptor nucleic acid can be produced based on the information provided herein along with standard recombinant nucleic acid techniques. Examples of references describing recombinant nucleic acid techniques include Molecular Cloning, Sambrook et al., Cold Spring Harbor Laboratory Press (1989); and Current Protocols in Molecular Biology, Frederick et al., John Wiley & Sons, Inc. (1995).

Due to the degeneracy of the genetic code different nucleic acid sequences can encode for a particular polypeptide. Thus, a large number of nucleic acids encoding for a receptor having the same amino acid sequence can be produced.

An embodiment of the present invention uses human nucleic acid encoding for the domains from CaR, GABABRIA, GABABRIB, GABABRI and/or mGluR8. The amino acid sequences of different domains is provided in Figures 1-3.

Recombinant Cells

Nucleic acid expressing a functional G-Protein fusion or a chimeric receptor can be used to create transfected cells lines expressing such receptors. Such cell lines have a variety of uses such as being used for high-throughput screening for compounds modulating receptor activity; being used to assay binding to the receptor; and as factories to produce large amounts of a receptor.

A variety of cell lines can couple exogenously expressed receptors to endogenous functional responses. Cell lines such as NIH-3T3, HeLa, NG115, CHO, HEK 293 and COS7 which are expected to lack CaR, mGluR8, and GABA_BR can be tested to confirm that they lack these receptors.

Production of stable transfectants can be accomplished by transfection of an appropriate cell line with, for example, an expression vector such as pMSG vector, in which the coding sequence for the G-protein fusion or chimeric GABA_ER cDNA has been cloned. Expression vectors containing a promoter region, such as

the mouse mammary tumor virus promoter (MMTV), drive high-level transcription of cDNAs in a variety of mammalian cells. In addition, these vectors contain genes for selecting cells stably expressing cDNA of interest. The selectable marker in the pMSG vectors encode an enzyme, xanthine-guanine phosphoribosyl transferase (XGPRT), conferring resistance to a metabolic inhibitor that is added to the culture to kill nontransfected

The most effective method for transfection of eukaryotic cell lines with plasmid DNA varies with the given cell type. The expression construct will be introduced into cultured cells by the appropriate technique, such as Ca² phosphate precipitation, DEAE-dextran transfection, lipofection or electroporation.

Expression of the receptor cDNA in cell lines can be assessed by sclution hybridization and Northern blot analysis.

cells.

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Binding Assavs

The present invention also includes using G-protein fusion receptors or chimeric GABA_BR in a binding assay. G-protein fusion receptors or chimeric GABA_BRs having a particular GABA_BR domain can be used, for example to facilitate obtaining compounds able to bind to that particular receptor domain; and to determine whether a compound which binds to a particular domain. For example, in a complete chimeric GABA_BR containing extracellular, transmembrane, and intracellular domains, the presence of one or more domains from CaR or mGluR are useful to present GABA_BR domain(s) to a binding agent in a form more like the GABA_BR domain(s) in the wild type receptor compared to an incomplete GABA_BR receptor fragment lacking one or more domains.

Binding assays can be carried out using techniques well known in the art. Binding assays preferably employ radiolabeled binding agents.

An example of a binding procedure is carried out by first attaching chimeric $GABA_BR$ to a solid-phase support to create an affinity matrix. The affinity matrix is then contacted with

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potential $GABA_{\mu}R$ binding agents. A large library of compounds may be used to determine those compounds binding to the affinity matrix. Bound compounds can be eluted from the column.

Therapeutic Modulation

As pointed out above, different types of diseases and disorders can be treated using compounds modulating CaR, mGluR, or GABA_BR activity. Additionally, such compounds can be used prophylactically. Compounds modulating GABA_BR2 activity can be administered to patients who would benefit from such treatment. Patients are mammals, preferably humans.

Modulators of CaR, mGluR, or GABA_BR activity can be administered to a patient using standard techniques. Techniques and formulations generally may be found in <u>Remington's Pharmaceutical Sciences</u>, 18th ed., Mack Publishing Co., Easton, PA, 1990 (hereby incorporated by reference herein).

Suitable dosage forms, in part, depend upon the use or the route of entry, for example, oral, transdermal, transmucosal, or by injection (parenteral). Such dosage forms should allow the therapeutic agent to reach a target cell whether the target cell is present in a multicellular host or in culture. For example, pharmacological compounds or compositions injected into the blood stream should be soluble. Other factors are well known in the art, and include considerations such as toxicity and dosage forms which retard the therapeutic agent from exerting its effect.

Therapeutic compounds can be formulated as pharmaceutically acceptable salts and complexes thereof. Pharmaceutically acceptable salts are non-toxic salts in the amounts and concentrations at which they are administered. The preparation of such salts can facilitate the pharmacological use by altering the physical characteristics of the compound without preventing it from exerting its physiological effect. Useful alterations in physical properties include lowering the melting point to facilitate transmucosal administration and increasing the

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solubility to facilitate administering higher concentrations of the drug.

The pharmaceutically acceptable salt of a compound may be present as a complex. Examples of complexes include an 8-chlorotheophylline complex (analogous to, e.g., dimenhydrinate:diphenhydramine 8-chlorotheophylline (1:1) complex; Dramamine) and various cyclodextrin inclusion complexes.

Pharmaceutically acceptable salts include acid addition salts such as those containing sulfate, hydrochloride, fumarate, maleate, phosphate, sulfamate, acetate, citrate, lactate, tartrate, methanesulfonate, ethanesulfonate, benzenesulfonate, ptoluenesulfonate, cyclohexylsulfamate and quinate.

pharmaceutically acceptable salts can be obtained from acids such as hydrochloric acid, maleic acid, sulfuric acid, phosphoric acid, sulfamic acid, acetic acid, citric acid, lactic acid, tartaric acid, malonic acid, methanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, cyclohexylsulfamic acid, fumaric acid, and quinic acid.

Pharmaceutically acceptable salts also include basic addition salts such as those containing benzathine, chloroprocaine, choline, diethanolamine, ethylenediamine, meglumine, procaine, aluminum, calcium, lithium, magnesium, potassium, sodium, ammonium, alkylamine, and zinc, when acidic functional groups, such as carboxylic acid or phenol are present. For example, see Remington's Pharmaceutical Sciences, 18th ed., Mack Publishing Co., Easton, PA, p. 1445, 1990. Such salts can be prepared using the appropriate corresponding bases.

Carriers or excipients can also be used to facilitate administration of therapeutic agents. Examples of carriers include calcium carbonate, calcium phosphate, various sugars such as lactose, glucose, or sucrose, or types of starch, cellulose derivatives, gelatin, vegetable oils, polyethylene glycols and physiologically compatible solvents. Examples of physiologically compatible solvents include sterile solutions of water for injection (WFI), saline solution and dextrose.

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GABA,R modulating compounds can be administered by different routes including intravenous, intraperitoneal, subcutaneous, intramuscular, oral, topical (transdermal), or transmucosal administration. For systemic administration, oral administration is preferred. For oral administration, for example, the compounds can be formulated into conventional oral dosage forms such as capsules, tablets, and liquid preparations such as syrups, elixirs, and concentrated drops.

Alternatively, injection (parenteral administration) may be used, e.g., intramuscular, intravenous, intraperitoneal, and 10 subcutaneous. For injection, compounds are formulated in liquid solutions, preferably, in physiologically compatible buffers or solutions, such as saline solution, Hank's solution, or Ringer's solution. In addition, the compounds may be formulated in solid form and redissolved or suspended immediately prior to use. 15 Lyophilized forms can also be produced.

Systemic administration can be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are well known in the art, and include, for example, for transmucosal administration, bile salts and fusidic acid derivatives. In addition, detergents may be used to facilitate permeation. Transmucosal administration, for example, may be through nasal 25 sprays, rectal suppositories, or vaginal suppositories.

For topical administration, compounds can be formulated into ointments, salves, gels, or creams, as is well known in the art.

The amounts of various GABABR modulating compounds to be administered can be determined by standard procedures taking into account factors such as the compound IC_{50} , EC_{50} , the biological half-life of the compound, the age, size and weight of the patient, and the disease or disorder associated with the patient. The importance of these and other factors to be considered are well known to those of ordinary skill in the art. Generally, the amount is expected to preferably be between about 0.01 and 50 mg/kg of the animal to be treated.

EXAMPLES

Examples are provided below illustrating different aspects and embodiments of the present invention. The examples include techniques that can be used to produce and use G-protein fusion-receptors and chimeric receptors. These examples are not intended to limit the claimed invention.

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Example 1: Construction of G-Protein Fusions

This example illustrates different G-protein fusion receptor constructs and techniques used to produce different G-protein fusion receptor constructs. Numbering of nucleotide position for all the following constructs is such that nucleotide number 1 corresponds to the A of the ATG start codon of the nucleotide sequence encoding the designated protein.

I. FULL-LENGTH CONSTRUCTS

20 A. phCaR

The cDNA encoding the human CaR (Garrett et al., (1995) J. Biol. Chem.270:12919) is harbored in the Bluescript SK(-) plasmid (Stratagene). This construct is referred to as phCaR.

25 B. phmGluR2

A full length human mGluR2 cDNA was amplified from human cerebellum MarathonReady cDNA (Clontech) using PCR primers based on the human mGluR2 cDNA sequence (Genbank Accession # 4504136). The obtained PCR fragment was subcloned into the pT7Blue TA vector (Novagen). A Hind III-Not I fragment containing the human mGluR2 cDNA was then subcloned into the Bluescript SKII(-) plasmid (Stratagene). This construct is referred to as phmGluR2.

C. phGaq

A full length human $G\alpha_q$ cDNA was amplified from human cerebral cortex Quick-Clone cDNA (Clontech) using PCR primers based on the human $G\alpha_q$ cDNA sequence (Genbank Accession # 4564044). The obtained PCR fragment was subcloned into the

Bluescript SKII(-) plasmid (Stratagene). This construct is referred to as $phG\alpha_{\bf q}$.

D. phmGluR8

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The cDNA encoding the full length human mGluR8 cDNA (Stormann et al., International Publication No. WO97/48724) is harbored in the Bluescript SKII(-) plasmid (Stratagene). This construct is referred to as phmGluR8.

 $10 II. G\alpha qi5$

The cDNA encoding the human $G\alpha_q$ i5 cDNA (Conklin et al (1993) Nature 363:274-77) is harbored in the Bluescript SKII(-) plasmid (Stratagene). This construct is referred to as $G\alpha_q$ i5. The nucleic acid and amino acid sequences for $G\alpha_q$ i5 are provided by SEQ. ID. NOs. 28 and 29 respectively.

III. phCaR/hmGluR2

This chimera contains the extracellular domain of the human CaR and transmembrane domain and intracellular cytoplasmic tail of human mGluR2. The chimeric junction between the CaR and hmGluR2 was created using a recombinant PCR strategy similar to those described above.

The first reaction used two primers, CAll56 (sense 19-mer, corresponding to nucleotides 1156-1174 of human CaR), and the hybrid primer CA/2 (antisense 42-mer, containing 21 nucleotides complementary to nucleotides 1774-1794 of human CaR and 21 nucleotides complementary to nucleotides 1660-1680 of the human mGluR2). These primers were used to amplify a 659 bp PCR fragment of human CaR.

In a separate PCR reaction using phmGluR2 as template, a 692 bp fragment of the human mGluR2 was amplified using a hybrid primer 2/CA (sense 42-mer, exactly complementary to primer CA/2) and oligo 2-2330m, (antisense 23-mer, complementary to nucleotides 2309-2331 of the human mGluR2 cDNA). The two PCR products generated from the above two reactions were annealed together in equimolar ratios in the presence of the external primers CA1156 and 2-2330m, and the Pfu DNA polymerase (Stratagene).

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The resulting chimeric PCR product was digested with SexAl (Boehringer Mannheim) and BamHI (New England Biolabs) and subcloned into phCaR digested with the same two restriction enzymes. In the final cloning step, the 3' end of human mGluR2 was subcloned into this construct using the restriction enzymes BsrGI and BamHI (both New England Biolabs). The sequence of the resultant chimeric construct, phCaR/hmGluR2, was verified by ABI automated DNA sequence analysis.

IV. phCaR/hmGluR2*Gqi5

This construct contains the phCaR/hmGluR2 chimeric receptor fused to human $G\alpha_{qi}5$. A HindIII-BamHI fragment containing the phCaR/hmGluR2 construct was subcloned into pcDNA3.1/Hygro(+) (Invitrogen) to aid in constructing this fusion protein. The chimeric junction between the C-terminus of phCaR/hmGluR2 and the N-terminus of $G\alpha_{qi}5$ was created using a recombinant PCR strategy similar to those described above.

The first reaction used two primers, 2-1713 (sense 21-mer, corresponding to nucleotides 1710-1730 of human mGluR2) and the hybrid primer 2/Q (antisense 42-mer, containing 21 nucleotides complementary to nucleotides 2596-2616 of human mGluR2, and 21 nucleotides complementary to nucleotides 1-21 of pG α qi5). These primers were used to amplify a 927 bp PCR fragment of phCaR/hmGluR2. In a separate PCR reaction all of G α qi5 was amplified using a hybrid primer Q/2 (sense 42-mer, exactly complementary to primer 2/Q) and the and the T3 primer commercially available from Stratagene.

These two PCR products generated from the above two reactions were annealed together in equimolar ratios in the presence of the external primers 2-1713 and T3, and the Pfu DNA polymerase (Stratagene). The resulting chimeric PCR product was digested with Bsu361 and BamHI (New England Biolabs) and subcloned into phCaR/hmGluR2 digested with the same two restriction enzymes. The sequence of the resultant chimeric fusion construct, phCaR/hmGluR2*G α qi5, was verified by DNA sequence analysis.

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V. phmGluR2//CaR Construct

This chimera contains the extracellular and transmembrane domains of human mGluR2 linked to the intracellular cytoplasmic tail domain of the human CaR. The chimeric junction was created using three separate PCR reactions.

The first reaction used two primers, 2-1713 (sense 21-mer, corresponding to nucleotides 1710-1730 of human mGluR2, Genbank Accession # 4504136) and the hybrid primer 2/CT (antisense 42-mer, containing 21 nucleotides complementary to nucleotides 2452 - 2472 of human mGluR2 and 21 nucleotides complementary to nucleotides 2602-2622 of the human CaR). These primers were used to amplify a 783 bp PCR fragment of human mGluR2. In a separate PCR reaction using phCaR in the BlueScript SK plasmid as template, a 750 bp fragment of the human CaR was amplified using a hybrid primer CT/2 (sense 42-mer, exactly complementary to primer 2/CT) and the T3 primer commercially available from Stratagene.

The two PCR products generated from the above two reactions were annealed together in equimolar ratios in the presence of the external primers 2-1713 and T3, and the Pfu DNA polymerase (Stratagene). The resulting chimeric PCR product was digested with BsrG I and Not I (New England Biolabs) and subcloned into pmGluR2 digested with the same two restriction enzymes. The sequence of the resultant chimeric construct, pmGluR2//CaR, was verified by ABI automated DNA sequence analysis.

VI. pmGluR2//CaR*Gαqi5 Construct

This construct contains the hmGluR2//CaR chimeric receptor fused to human $G\alpha_{\bf q}{\rm i5}$. The chimeric junction between the C-terminus of hmGluR2//CaR and the N-terminus of $G\alpha_{\bf q}{\rm i5}$ was created using a recombinant PCR strategy similar to that described above for the construction of phmGluR2//CaR.

The first reaction used two primers, CRP10A (sense 18-mer, corresponding to nucleotides 2812-2829 of phCaR) and the hybrid primer CaRQ (antisense 42-mer, containing 21 nucleotides complementary to nucleotides 3214- 3234 phCaR, and 21 nucleotides complementary to nucleotides 1-21 of pG α_q i5). These primers were used to amplify a 443 bp PCR fragment of hmGluR2//CaR. In a

separate PCR reaction, all of $G\alpha_{Q1}5$ was amplified using a hybrid primer QCaR (sense 42-mer, exactly complementary to primer CaRQ) and the T3 primer commercially available from Stratagene.

The two PCR products generated from the above two reactions were annealed together in equimolar ratios in the presence of the external primers CRP10A and T3, and the Pfu DNA polymerase (Stratagene). The resulting chimeric PCR product was digested with BstE II and Not I (New England Biolabs) and subcloned into pmGluR2//CaR digested with the same two restriction enzymes. The sequence of the resultant chimeric fusion construct, pmGluR2//CaR*G α_{q} i5, was verified by ABI automated DNA sequence analysis.

VII. Fusion Receptor Protein Linker Addition Constructs

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A. phmGluR2//CaR*AAA*Gaai5

A linker encoding three alanine residues was incorporated into the phmGluR2//CaR*G α_q i5 construct by mutagenesis (Stratagene QuickChange Mutagenesis Kit). A sense 40-mer, 2CQ+LP, contained 16 nucleotides corresponding to 3219-3234 of human CaR, followed by the 9 nucleotide sequence (GCGGCCGCC) encoding three alanine residues and a NotI restriction enzyme site, and then 15 nucleotides corresponding to nucleotides 1-15 of G α_q i5. 2CQ+LP was annealed to an antisense 40-mer, 2CQ+LM, the exact complement of 2CQ+LP. These oligos were used in the mutagenesis reaction according to the manufacturer's protocol. Restriction enzyme analysis and DNA sequence analysis confirmed the insertion of the 9 nucleotide linker (GCGGCCGCC) between the C-terminus of phmGluR2//CaR and the N-terminus of G α_q i5. This construct was designated phmGluR2//CaR*AAA*G α_q i5.

B. Human GABA_BR2*AAA*Gα₉c5 and human GABA_BRla*AAA*Gα₉c5

These constructs contain the human GABA_BR2 (hGABA_BR2: Genbank Accession # AJ 012188) and human GABA_BR1a (hGABA_BR1a: Genbank Accession # AJ 012185) fused at their C-terminus to the N-terminus of human G α_q o5 (hG α_q o5: Nature 363:274-276, 1993). Human GABA_BR2, hGABA_BR1a, and hG α_q o5 were cloned into the plasmid pcDNA3.1/Hygro+ (Invitrogen) and are designated phGABA_BR2,

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phGABA_BR1a, and phG α_q o5. The first reaction used two primers, XcmI-R2 (sense 20-mer, corresponding to nucleotides 2650-2669 of phGABA_BR2) and the hybrid primer R2/Go5(-) (antisense 45-mer, containing 18 nucleotides complementary to nucleotides 2806-2823 of phGABA_BR2 and 18 nucleotides complementary to nucleotides 1-18 of hG α_q o5). These two complementary areas flank a 9 nucleotide sequence coding for 3 alanine sequences with a unique NotI restriction site. These primers were used to amplify a 200 base-pair PCR fragment.

In a separate PCR reaction, part of $hG\alpha_qo5$ was amplified using a hybrid primer $R2/G\alpha_qo5(+)$ (sense 45-mer), exactly complementary to R2/Go5(-) and XbaI-Go5 primer (22-mer containing 22 nucleotides complementary to nucleotides 873-895 of $hG\alpha_qo5$) These primers were used to amplify a 914 base-pair PCR product. The two PCR products generated from the above two reactions were annealed together in equimolar ratios in the presence of the external primers; XcmI-R2 and XbaI-Go5, and Pfu polymerase (Stratagene).

The resulting chimeric PCR product was digested with the restriction endonucleases XcmI and XbaI (New England Biolabs) and subcloned into phGABA_BR2 digested with the same two restriction enzymes. The resulting clone was then digested with HindIII and XbaI and subcloned into phG α_q o5 cut with HindIII and XbaI resulting in the chimeric hGABA_BR*AAA*G α_q o5. The chimeric junction between the C-terminus hGABA_BRla, the Ala linker, and the N-terminus cf hG α_q o5 was created using a recombinant PCR strategy similar to those described above.

To construct hGABA_BRla*AAA*Gqo5, the first reaction used a commercially available T7 primer (Novagen) and the NtI hGBRl primer (CAGAGTCATGGCGGCCGCCTTATAAAGCAAATGCACTCG) corresponding to nucleotide numbers 1-9 of hG α_q o5 and nucleotide numbers 2863-2883 of hGABA_PRla.

VIII. phmGluR8//CaR Construct

This chimera contains the extracellular and transmembrane domains of human mGluR8 linked to the intracellular cytoplasmic tail domain of the human CaR. The chimeric junction between hmGluR8 and the CaR was created using a recombinant PCR strategy

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similar to those described above.

The first reaction used two primers, CH5A (sense 19-mer, corresponding to nucleotides 2187-2205 of human mGluR8, Stormann et al., International Publication Number WO97/48724) and the hybrid primer CH5B (antisense 40-mer, containing 22 nucleotides complementary to nucleotides 2523 - 2544 of human mGluR8, and 18 nucleotides complementary to nucleotides 2602-2619 of the human CaR). These primers were used to amplify a 375 bp PCR fragment of human mGluR8. In a separate PCR reaction using phCaR in the BlueScript SK(-) plasmid as template, a 750 bp fragment of the human CaR was amplified using a hybrid primer CH5C (sense 40-mer, exactly complementary to primer CH5B) and the T3 primer commercially available from Stratagene.

The two PCR products generated from the above two reactions were annealed together in equimolar ratios in the presence of the external primers CH5A and T3, and the Pfu DNA polymerase (Stratagene). The resulting chimeric PCR product was digested with BsrG I and Xba I (New-England Biolabs) and subcloned into pmGluR8 digested with the same two restriction enzymes. The sequence of the resultant chimeric construct, pmGluR8//CaR, was verified by DNA sequence analysis.

IX. mGluR8//CaR*Gαqi5 Construct

This construct contains the hmGluR8//CaR chimeric receptor fused to human G α_q i5. The chimeric junction between the C-terminus of hmGluR8//CaR and the N-terminus of G α_q i5 was created using a recombinant PCR strategy similar to that described above for the construction of phmGluR2//CaR*G α_q i5.

The first reaction used two primers, CRP10A (sense 18-mer, corresponding to nucleotides 2812-2829 of phCaR) and the hybrid primer Gqi5/CaR (antisense 40-mer, containing 21 nucleotides complementary to nucleotides 3214-3234 phCaR, and 19 nucleotides complementary to nucleotides 1-19 of pG $\alpha_{\rm q}$ i5). These primers were used to amplify a 441 bp PCR fragment of hmGluRB//CaR.

In a separate PCR reaction all of $G\alpha_{Q}i5$ was amplified using a hybrid primer CaR/Gqi5 (sense 40-mer, exactly complementary to primer Gqi5/CaR) and the Apa I-mut primer (20-mer). The two PCR products generated from the above two reactions were annealed

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together in equimolar ratios in the presence of the external primers CRP10A and Apa I-mut, and the Pfu DNA polymerase (Stratagene).

The resulting chimeric PCR product was digested with BstE II and Apa I (New England Biolabs) and subcloned into pmGluR8//CaR digested with the same two restriction enzymes. The sequence of the resultant chimeric fusion construct, pmGluR8//CaR*G α_q i5, was verified by DNA sequence analysis.

10 Example 2: Functional Expression of CaR/GABA_BR2

In vitro transcribed RNA (7 ng) encoding a chimeric $CaR/GABA_BR2 \ (CaR \ extracellular \ and \ transmembrane \ domains, \ and intracellular \ GABA_BR2 \ domain) \ was \ co-injected \ with \ in \ vitro transcribed RNA (2 ng) encoding <math display="inline">G_{\alpha}15$ into Xenopus oocytes.

15 Following a 72-hour incubation, the oocytes were voltage-clamped using standard electrophysiological techniques (Hille, B., <u>Ionic Channels of Exictable Membranes</u>, pp.30-33, Sinauer Associates, Inc., Sunderland, Ma., 1992). Activation of the chimeric receptor was detected by increases in the calcium-activated chloride current.

Application of the CaR activator 100 μ M Gd³*, resulted in reversible, oscillatory increases in the calcium-activated chloride current as shown in Figure 8. These data demonstrate the functional response of the chimeric CaR/GABA_R2 receptor upon activation via a site within the CaR extracellular domain. In this assay, the G $_{\alpha}$ 15 subunit acts to promote signal transduction through intracellular pathways that mobilize intracellular Ca**.

Example 3: Expression of Different G-Protein Fusion Receptors

30 The ability of different G-protein fusions to transduce signal resulting from ligand binding is shown in Figure 15. The different G-protein fusion receptors used in this example were as follows: mGluR2//CaR'Gqi5 (SEQ. ID. NO. 37), CaR/mGluR2'Gqi5 (SEQ. ID. NO. 33), and mGluR8//CaR'Gqi5 SEQ. ID. NO. 41.

Occytes suitable for injection were obtained from adult female Xenopus laevis toads using procedures described in C. J.

Marcus-Sekura and M. J. M. Hitchcock, Methods in Enzymology, Vol. 152 (1987).

Receptor fusion cRNAs were dissolved in water and 50 nl (12.5 ng/oocyte) were injected into individual oocytes.

5 Following injection, occytes were incubated at 16°C in MBS containing 1 mM $CaCl_2$ for 2 to 7 days prior to electrophysiological recording.

Test substances were applied by superfusion at a flow rate of about 5 ml/min. Receptor fusion activation was determined by measuring the increase in calcium-activated chloride current (I_{cl}). Increases in I_{Cl} were quantified by measuring the peak inward current stimulated by activating agent, relative to the holding current at -60 mV. Application of 100 µM L-glutamate elicited a response from the mGluR2//CaR*Gaqi5 and mGluR8//CaR*Gaqi5. Application of 100 µM Gd³* activated the CaR/mGluR2 Gqi5.

Other embodiments are within the following claims. Thus, while several embodiments have been shown and described, various modifications may be made, without departing from the spirit and scope of the present invention.

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Claims

A G-protein fusion receptor comprising
 an extracellular domain comprising an extracellular domain

 amino acid sequence substantially similar to either an

amino acid sequence substantially similar to either an extracellular CaR amino acid sequence, an extracellular mGluR amino acid sequence, or an extracellular GABA_B receptor amino acid sequence;

a transmembrane domain joined to the carboxy terminus of said extracellular domain, said transmembrane domain comprising a transmembrane domain amino acid sequence substantially similar to either a transmembrane CaR amino acid sequence, a transmembrane mGluR amino acid sequence, or a transmembrane GABAs receptor amino acid sequence;

an intracellular domain joined to the carboxy terminus of said transmembrane domain comprising all or a portion of an intracellular amino acid sequence substantially similar to either an intracellular CaR amino acid sequence, an intracellular mGluR amino acid sequence, or an intracellular GABAB receptor amino acid sequence, provided that said portion is at least about 10 amino acids;

an optionally present linker joined to the carboxy terminus of said intracellular domain; and

a G-protein joined either to said intracellular domain or to said optionally present linker, provided that said G-protein is joined to said optionally present linker when said optionally present linker is present.

- 2. The G-protein fusion receptor of claim 1, wherein said extracellular domain consists of said extracellular domain amino acid sequence, said transmembrane domain consists of said transmembrane domain amino acid sequence; and said intracellular domain consists of said transmembrane domain amino acid sequence.
- 35 3. The G-protein fusion receptor of claim 2, wherein said optionally present linker is present and is a polypeptide 3 to 30 amino acids in length.

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- 4. The G-protein fusion receptor of claim 2, wherein said optionally present linker is not present.
- 5. The G-protein fusion receptor of claim 3 or 4, wherein said G-protein is selected from the group consisting of: $G_{\alpha_{15}}$. $G_{\alpha_{16}}$, Gqo5, and Gqi5
- 6. The G-protein fusion of claim 5, wherein any of said CaR sequence present is a human CaR sequence, any of said mGluR sequence present is from a human mGluR, and any of said GABA_B receptor sequence present is from human mGluR.
 - 7. A nucleic acid comprising a nucleotide sequence encoding for the G-protein fusion of any one of claims 1-6.

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- 8. An expression vector comprising a nucleotide sequence encoding for the G-protein fusion of any one of claims 1-6 transcriptionally coupled to a promoter.
- 9. A recombinant cell comprising the expression vector of claim 8 and a cell wherein the G-protein fusion is expressed and is functional.
- 10. A recombinant cell produced by combining a vector comprising the nucleic acid of claim 9 and elements for introducing heterologous nucleic acid into a cell wherein the G-protein fusion receptor is expressed, and said cell.
- 11. A process for the production of a G-protein fusion 30 receptor comprising:

growing procaryotic or eukaryotic host cells comprising a nucleic acid sequence expressing the G-protein fusion receptor of any one of claims 1-6, under suitable nutrient conditions allowing for cell growth.

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- 12. A method of measuring the ability of a compound to effect G-protein fusion activity comprising the steps of:
- a) providing said compound to a cell expressing the G-protein fusion receptor of any one of claims 1-6, and

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- b) measuring the ability of said compound to affect the activity of said receptor as an indication of the ability of said compound to effect G-protein fusion receptor activity.
- 5 13. A chimeric receptor comprising

an extracellular domain comprising an extracellular domain amino acid sequence substantially similar to a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4 and SEQ ID NO: 5;

a transmembrane domain comprising a transmembrane domain amino acid sequence substantially similar to a sequence selected from the group consisting of SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, and SEQ ID NO: 10; and

an intracellular cytoplasmic domain comprising an intracellular domain amino acid sequence substantially similar to a sequence selected from the group consisting of SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, and SEQ ID NO: 14;

wherein at least one domain is present which comprises an amino acid sequence substantially similar to a sequence selected from the group consisting of: SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 7, SEQ ID NO: 8, and SEQ ID NO: 9, SEQ ID NO: 12, SEQ ID NO: 13, and SEQ ID NO: 14; and at least one domain is present which comprises an amino acid sequence substantially similar to a sequence selected from the group consisting of: SEQ ID NO: 1, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 10, SEQ ID NO: 11, and SEQ ID NO: 15.

- 14. The chimeric receptor of claim 13 wherein said extracellular domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID NOS: 2, 3, and 4; said transmembrane domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID Nos: 6, 7, 8, 9, and 10; and said intracellular domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID Nos: 11 and 15.
- 15. The chimeric receptor of claim 14, wherein said extracellular domain has a sequence similarity of at least 90%

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with the amino acid sequence of SEQ ID NO: 2; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence SEQ ID NO: 7; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEO ID NO: 11.

- 16. The chimeric receptor of claim 14, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 3; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 8; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 11.
- 17. The chimeric receptor of claim 14, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence SEQ ID NO: 4; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 9; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO 11.
- extracellular domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID Nos: 1, 2, 3, 4 and 5; said transmembrane domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID Nos: 7, 8, and 9; and said intracellular domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID Nos: 11, 12, 13, 14, and 15.
- 19. The chimeric receptor of claim 18, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 7; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 11.

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- 20. The chimeric receptor of claim 18, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 8; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO 11.
- 21. The chimeric receptor of claim 18, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 9; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO 11.
- 22. The chimeric receptor of claim 13, wherein said extracellular domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID NOs: 1, 2, 3, 4, and 5; said transmembrane domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID Nos: 6, 7, 8, 9, and 10; and said intracellular domain has a sequence similarity of at least 90% with an amino acid sequence selected from the group consisting of SEQ ID NOs: 12, 13, and 14.
 - 23. The chimeric receptor of claim 22, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 6; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 12.
 - 24. The chimeric receptor of claim 22, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino

acid sequence of SEQ ID NO: 7; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 12.

- 25. The chimeric receptor of claim 22, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 8; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 13.
 - 26. The chimeric receptor of claim 22, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 6; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 13.

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- 27. The chimeric receptor of claim 22, wherein said extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 9; and said intracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 14.
- 28. The chimeric receptor of claim 22, wherein said

 30 extracellular domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 1; said transmembrane domain has a sequence similarity of at least 90% with the amino acid sequence of SEQ ID NO: 6; and said intracellular domain has a sequence similarity of at least 90% with the amino acid

 35 sequence of SEQ ID NO: 14.
 - 29. The chimeric receptor of any one of claims 13-28, wherein said receptor functional couples to a G-protein.

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- 30. The chimeric receptor of any one of claims 13-28, wherein said chimeric receptor consists of said extracelluar domain, said transmembrane domain, said intracellular domain, and an optionally present G-protein α subunit covalently joined to said intracellular domain.
- 31. The chimeric receptor of claim 30, wherein said chimeric receptor consists of said extracelluar domain, said transmembrane domain, and said intracellular domain.
- 10 32. The chimeric receptor of claim 30, wherein said G-protein α subunit consists of the amino acid sequence of SEQ ID Nos: 16 or 17.
- 33. A nucleic acid comprising a nucleotide sequence encoding for the chimeric receptor of any one of claims 13-32.
- 34. An expression vector comprising a nucleotide sequence encoding for the chimeric receptor of any one of claims 13-32 transcriptionally coupled to a promoter.
 - 35. A recombinant cell comprising the expression vector of claim 34 and a cell wherein the chimeric receptor is expressed and is functional.
 - 36. A recombinant cell produced by combining a vector comprising the nucleic acid of claim 33 and elements for introducing heterologous nucleic acid into a cell wherein the chimeric receptor is expressed, and said cell.
 - 37. A process for the production of a chimeric receptor comprising:

growing procaryotic or eukaryotic host cells comprising a nucleic acid sequence expressing the chimeric receptor of any one of claims 13-32, under suitable nutrient conditions allowing for cell growth.

38. A method of measuring the ability of a compound to effect $GABA_BR$ or mGluR activity comprising the steps of:

- a) providing said compound to a cell expressing the chimeric receptor of any one of claims 13-32, and
- b) measuring the ability of said compound to affect the activity of said receptor as an indication of the ability of said compound to effect GABA $_{\rm B}R$ or mGluR activity.
 - 39. The method of claim 38, wherein said method measures activity at a $GABA_{\mu}R$.
- 40. The method of claim 38, wherein said method measures activity at a mGluR.
- 41. A fusion receptor polypeptide comprising a receptor and a G-protein α subunit, wherein said G-protein α subunit is fused to the intracellular domain of said receptor, provided that said receptor is not an adrenoreceptor.

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	SEQ	רוו	. 4	λ/	Ι Δ	Ŧ	Y	ç	\mathcal{C}	_	w	ν	T	t	Δ	Ť	т	·u	, LI	7	,	Δ	v	c.	Đ	n	. ^	
	SEQ SEQ SEQ	ID ID ID	2 3 4	M M	I L I G	L P S	L G P E	L A R	L P S	L F S	A A G	P R Q	L V P	F G	L W P	R P X	P L P	P P P	G L P	A L P	G V P	G V P	A M P	Q A A	T A R	P G L	N V L	
	SEQ SEQ SEQ SEQ	ID ID ID	2 3 4	T P L	S V L	E W L	K G A L	C S P	Q H L	I S L	I P L	H H P	P L L	P P A	W R P	E P G	G H A	G S W	I R G	R V W	Y P A	R P R	G H G	L P A	T S P	R S R	D E P	
9	SEQ SEQ SEQ SEQ	ID ID ID	2 3 4	V R P	K A S	A V S	R I Y P V	N I P	F G L	L A S	P L I	V F M	D P G	Y M L	E S M	I G P	E G L	Y W T	V P K	C G E	R G V	G Q A	E A K	R C G	E Q S	V P I	V A G	
S	SEQ SEQ SEQ SEQ SEQ	ID ID ID	2 3 4	P E G	K M V	V A L	E R L P	K E A	C D V	L V E	A N L	N S A	G R I	S R E	W D Q	T I I	D L R	M P N	D D E	T Y S	P E L	S L L	R K R	C L P	V I Y	R H F	I H L	1
5	SEQ SEQ SEQ SEQ	ID ID ID	2 3 4	S S L	K K R	S C L	V Y D Y D	L P D	T G T	L Q E	E A C	N T D	G K N	K Y A	V L K	F Y G	L E L	T L K	G L A	G Y F	D N Y	L D D	P P A	A I I	L K K	D I Y	G I G	1
5	SEQ SEQ SEQ SEQ SEQ	ID ID ID	2 3 4	R M N	V P H	D G L	М	R S V	C S F	D V G	P S G	D T V	F L C	H V P	L A S	V E V	G A T	S A S	S R I	R M I	S W A	I N E	C L S	S I L	Q V Q	G L G	Q S W	`
S	EQ EQ EQ EQ	ID ID ID	2 3 4	S G L	T S V	P S Q	K S L	P P S	H A F	C L A	Q S A	> 7 7 T	N R T	R Q P	T R V	P F L	H P A	S T D	E F K	R F K	R R K	A T Y	V H P	Y P Y	I S F	G A F	A T R	

FIG. Ia.

SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	F P M S H N P T V P S D	G G W P R V K L N A V N	GGQ2 FEKV PAll	ACQPA WGWKK LKLLK	H Q A T A M A D 1 V E M A L E D V N 1 A T 1 Q Q T T E H Y Q W K R V G T F S R V V P P D S
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	SRRD VFTS LTQD	I L P D T L D D V Q R F	Y E L I L E E I S E V I	K L I H H : R V K E A : R N D L T :	D Y G R P G I E K D S K C D P G Q A G I E I T F R Q S G V L Y G E D I E Y V S T L A S E G
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	T K Y L F F S D I S D T	Y E L L P A V P E S F S	Y N D ! V K N I N D P (PIKIII LKRQD CTSVKI	L I S Q Y S D E E L M P G C S S V S A R I I V G L F Y K L K G N D V R I I G G V C I A Q S
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	T L V A E T E A 1 L G Q	E A A R R K V F F D Q N	M W N I C E V I M A A I	LIVLS YKERL: KVFCC:	I V V F S S G P D Y G S S S P A L S F G K K Y V W F L A Y E E N M Y G S K R L L E T P N A
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	NRQR 1GWY KYQW	FPTF ADNW IIPG	FRTH FKI WYE!	H P S A T Y D P S I I P S W W E	I W L A S E A W A L H N P T R V K L N C T V D E M T E Q V H T E A N S S E A A K L N Q S
SEQ ID 2 SEQ ID 3 SEQ ID 4	F E K W A V E G R C L R	G W K K H I T T K N L L	IAT EIV AAM	IQQTT MLNPA EGYIG	T 1 G F A L K A G E V F T S T L D D N T R S 1 S N M T V D F E P L S S K P V Y Q Q E E 1 A
SEQ ID 2 SEQ ID 3 SEQ ID 4	LEER SQEF QIKT	V K E A V E K L I S G K	GIE TKR TPQ	I T F R Q L K R H P Q Y E R E	S V H N G F A K E S F F S D P A V P E E T G G F Q E A Y N N K R S G V G D R Y F R S R T L

FIG. 1b.

SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	V K N P L A P S K	ETILKI YD2 FH(R Q D A I W 5 Y A) A : / A . Y :	R 1 L A D G	I V L A I W	G . L . V	L F N K I A	Y T K	E S T	T G L	E G Q	A G R	R G A	K R M	V S E	F G T
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	C E V V R L L H A	ESC YKI EDI SSF	ERL FNY RHQ	F (N)	G K N Q I Q	K Y T I D F	V T N	W F D Q Y T	I D	I Y H	G` R T	W I A I L (Y M G	A N R	D S I	N S I	W S L
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	F K I F E G N A M	V E T Y D I V S C I N E T	9 5 I 5 H V 7 N F	N O	C T F D G V	V D A S T G	E : G Q	M T S R V V	E M F	A A R	V W N	E (T : G :	G L E	H I R	I E M	T Q G	T L T
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	E I V Q G G I K F	HAI MLI SYI TQI	N P A	G S	T R Y Y R E	S 1 D S V K	S T V	N M K D G E	T D Y	S L N	Q S ' A	E : W : V :	F S A	V K D	E T T	K D L	L K E
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	TKR WIC	KVI LKI GGS! IDT:	RHP PPA IRF	E D (E T Q T G S	G G L V E P	F I P	Q E K T K D	A F K	P R T	L . F	A ' L I	Y S L	D Q E	A K Q	I L	W R
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	ALA	LAI	LNK	Τ	S G	G G	G	R S	G	V	R	L	E	D	F	N	Y
SEQ ID 1 SEQ ID 2 SEQ ID 3 SEQ ID 4 SEQ ID 5	NNC) TI'	IDÇ) I	Y R	A M	1 N	S S	S	F	Ē	G	V	S	G	H	ν

FIG. Ic.

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SEQID 1 EKILWSGFSREVPFSNCSRDCLAGT
SEQID 2 V F D A S G S R M A W T L I E Q L Q G G S Y K K I
SEQ ID 3
SEQ ID 4
SEQID 5 KKTVKGVPCCWHCERCEGYNYQVDE
SEQID 1 RKGII EGEPTCCFECVECPDGEYSD
SEQID 2 GYYDSTKDDLSWSKTDKWIGGSPPA
SEQ ID 3
SEQ ID 4
SEQID 5 L S C E L C P L D Q R P N M N R T G C Q L I P I I
SEQID 1 ETDASACNKCPDDFWSNENHTSCIA
SEQID 2 DQTLVIKTFRFLSQK
SEQ ID 3
SEQ ID 4
SEQID 5 KLEWHSPW
SEQID 1 KEIEFLSWTEPF
SEQ ID 2
SEQ ID 3
SEQ ID 4
SEQ ID 5
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FIG. Id.

SEO ID 6 SEO ID 7 SEO ID 8 SEO ID 9 SEO ID 10	LFIS LFIS LYSI	S V S V I S V S V I I L S A	S S L S S L T L	G	A V V C L S A V V C L S A S A F L F	F
SEQ ID 6 SEQ ID 7 SEQ ID 8 SEQ ID 9 SEQ ID 10	Y 1 Q N	N S Q P I N S Q P I M S S P '	1 L N N 1 L N N 7 M N N	L T A V L T A V L I I L	G C S L A L G C S L A L G G M L S Y	S S L F F I G E P A A V F P L G L D A A V F P L G L D A S I F L F G L D I T F L M I A A P
SEQ ID 6 SEQ ID 7 SEQ ID 8 SEQ ID 9 SEQ ID 10	G Y H I G Y H I G S F V	IGRN(IGRN(/SEK	OFPF OFPF FFET	V C Q A V C Q A L C T V	R L W L L G R L W L L G R T W I L T	C I L V K T N R V L G F S L G Y G S L G F S L G Y G S V G Y T T A F G A A L L T K T N R I
SEO ID 6 SEO ID 7 SEO ID 8 SEO ID 9 SEO ID 10	M F T K M F T K M F A K	(I W W Y (I W W Y (T W R Y	/	FTKK FTKK FKNV	E E K K E W K M K K K I	Q F L L V F L C T R K T L E P W K L R K T L E P W K L I K D Q K L L V I Q L V I T F S L I
SEQ ID 6 SEQ ID 7 SEQ ID 8 SEQ ID 9 SEQ ID 10	YATV YATV VGGM	/ G L L \ / G L L \ / L L I I	GMD GMD CLC1	V L T L V L T L L I C W	A I W Q I V	Q E L E D E 1 1 F D P L H R T 1 E T D P L H R T 1 E T R R T V E K Y S M Y G E Q R T L D P
SEO ID 6 SEQ ID 7 SEQ ID 8 SEQ ID 9 SEQ ID 10	F A K E F A K E E P D F	E E P K I E E P K I P A G R I		V S I L V S I L R P L L	P Q L E H C P Q L E H C E H C E N T	A A I C F F F A F S S R K M N T W L S S R K M N T W L H M T I W L G I V Y S I L L M V T C
SEQ ID 6 SEQ ID 7 SEQ ID 8 SEQ ID 9 SEQ ID 10	G F ' G F ' Y A Y I	Y G Y K Y G L L	G L L L G L L L M L F G	L L G I L L G I C F L A	I F L A Y E T I F L A Y E T W E T R N V	F F I V W I S F I K S V S T E K I N K S V S T E K I N S I P A L N D S K F T M Y T T C I I

SEQ ID 6 SEQ ID 7 SEQ ID 8 SEQ ID 9 SEQ ID 10	P. A. D. H. D. D. H. Y. W. I.	1 F 1 F	A A A A A A	V V ! S	G G V	М М Ү	AAN	1 1 V	Y Y G	N N 1	У У М	A A C	V V 1	L L	\circ	LLA	1 1 A	TTV	A A S	5 P F	V V L		M M R	1 1 D	.1 .1 0	SSP	S S Z	00>	000
SEQ ID 6 SEQ ID 7 SEQ ID 8 SEQ ID 9 SEQ ID 10	FDDFL	A A A A	F	A	F	A A V	S S	L	A A F	1 C	v s	F	S	S T	Y L	1 C	7	L	٧	٧	L	F	٧						

FIG. 2b.

```
SEQID 11 KPSRNTIE EVRCSTAAHAFKVAARATLRRS
SEQID 12 RRLITRGEWQSEAQDTMKTGSSTNNNEEEK
SEOID 13 RRLITRGEWOSEAODTMKTGSSTNNNEEEK
     ITLRTNPDAATONRRFOFTONOKKEDSKTS
SEQ ID 14
SEQID 15 HPEQNVQKRKRSFKAVVTAATMQSKLIQKG
SEOID 11 N V S R K R S S S L G G S T G S T P S S S I S S K S N S E D
SEQID 12 SALLEKENRELEKI I AEKEERVSELAHOLO
SEOID 13 SRLLEKENRELEKII AEKEERVSELRHQLQ
SEQID 15 NORPNGEVKSELCESLETNSKSSVE FPMVK
SEQID 11 P F P Q P E R Q K Q Q Q P L A L T Q Q E Q Q Q P L T L P Q
SEQID 12 SRQQLRSRRHPFTPPEPSGGLPRGPPEPPD
SEQID 13 SAQQLAS ARHPPTPPEPSGGLPAGPPEPPD
SEOID 14 L D K D L E E V T M Q L Q D T P E K T T Y I K Q N H Y Q E L
SEQID 15 SGSTS
SEQID 11 QQRSQQQPRCKQKVIFGSGTVTFSLSFDEP
SEQID 12 RLSCDGSRVHLLYK
SEOID 13 RLSCDGSRVHLLYK
SEQID 14 ND I LN L G N F T E S T D G G K A I L K N H L D Q N P Q L
SEQ ID 15
SEQID 11 QKNAMAH GNSTHQNSLEAQKSSDTLTRHQP
SEQ ID 12
SEQ ID 13
SEQID 14 QWNTTEPSRTCKDPIED INSPEHIQRRLSL
SEQ ID 15
SEQID 11 LLPLQCGETDLDLTVQETGLQGPVGGDQRP
SEQ ID 12
SEQ ID 13
SEQID 14 QLP I LHHAYLPS I GGVDASCVSPCVSPTAS
SEQ ID 15
SEOID 11 EVEDPEELS PALV VSSSQS FV ISGGGS TV T
SEQ ID 12
SEQ ID 13
SEOID 14 PRHRHVPPS FRVMVSGL
SEQ ID 15
```

FIG. 3a.

SEQ ID 11 E N V V N S SEQ ID 12 SEQ ID 13 SEQ ID 14 SEQ ID 15

FIG. 3b.

SEQ. ID. NO. 16 MARSLTWGCCPWCLTEEEKTAARIDQEINR SEQ. ID. NO. 17 MARSLTWRCCPWCLTEDEKAAARVDQEINB SEQ. ID. NO. 16 ILLEQKKQEREELKLLLLGPGESGKSTFIK SEQ. ID. NO. 17 ILLEQKKQDRGELKLLLLGPGESGKSTFIK SEQ. ID. NO. 16 QMR!!HGVGYSEEDRRAFRLL!YQN!FVSM SEQ. ID. NO. 17 QMRIIHGAGYSEEERKGFRPLVYQNIFVSM SEQ. ID. NO. 16 QAM! DAMDRLQIPFSRPDSKQHASLVMTQD SEQ. ID. NO. 17 RAMIEAMERLQIPFSRPESKHHASLVMSQD SEQ. ID. NO. 16 PYKVSTFEKPYAVAMQYLWRDAGIRACYER SEQ. ID. NO. 17 PYKVTTFEKRYAAAMQWLWRDAGIRACYER SEQ. ID. NO. 16 RREFHLLDSAVYYLSHLERISEDSYIPTAQ SEQ. ID. NO. 17 RREFHLLDSAVYYLSHLERITEEGYVPTAQ SEQ. ID. NO. 16 DVLRSRMPTTGINEYCFSVKKTKLRIVDVG SEQ. ID. NO. 17 DVLRSRMPTTGINEYCFSVQKTNLRIVDVG SEQ. ID. NO. 16 G Q R S E R R K W I H C F E N V I A L I Y L A S L S E Y D Q SEQ. ID. NO. 17 G Q K S E R K K W I H C F E N V I A L I Y L A S L S E Y D Q SEQ. ID. NO. 16 CLEENDQENRMEESLALFSTILELPWFKST SEQ. ID. NO. 17 CLEENNQENRMKESLALFGTILELPWFKST SEQ. 1D. NO. 16 SVILFLNKTDILEDKIHTSHLATYFPSFQG SEQ. ID. NO. 17 S V I L F L N K T D I L E E K I P T S H L A T Y F P S F Q G SEQ. ID. NO. 16 PRRDAEAAKS FILD M. YARV YAS CAEPQDGG SEQ. ID. NO. 17 PKQDAEAAKRFILDMYTRMYTGCVDGPEGS SEQ. ID. NO. 16 RKGSRARRFFAHFTCATDTQSVRSVFKDVR *SEQ. ID. NO. 17* KKGARSRRLFSHYTCATDTQN | RKVFKDVR

FIG. 4a.

SEO. ID. NO. 16 DSVLARYLDEINLL SEO. ID. NO. 17 DSVLARYLDEINLL

FIG. 4b.

WO 99/51641 PCT/US99/07333

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ClustalW Formatted Alignments

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A A	T T	G G	T G	T G	G G	C C	T C	G C	C G	T G	G G	C G	T C	G C	C	T C	A T	C T	T T	G T	G G	C C	G G C G	C
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C G	A G	C G	T T	C G	T G	T G	C G	C T	T G	C G	C C	G C	C A	C C	C T	C G	C C	C C	G G	G C	G T	C T	A G C C	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G G	G G	G T	C T	G G	G T	G G	G A	C T	G G	C G	A C	G G	A G	C	C A	C G	C G	C G	A G	A T	C G	G G	G C C T	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A C	C C	C G	T G	C T	A G	G T	A G	A G	G G	G C	T C	T T	G C	C	C C	A A	G C	A T	T C	C C	A C	T	C A C G	C C
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A A	C T	C C	C T	G C	C C	C C	C G	T C	G G	G G	G C	A C	A T	G C	G A	G C	G T	G C	C G	A C	T G	C G	T A G T	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C	T C	A C	C C	C C	G C	G G	G C	G A	C C	C C	T C	C	A T	C C	T C	C T	G C	G A	G G	A A	C A	C C	A A G C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G C	T G	G C	A G	A C	G A	G G	C T	T G	A T	T A	C	A A	A T	C C	T G	T G	C G	C G	T C	G A	C C	C T	A G	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T T	G T	G C	A C	C C	T A	A T	T G	G A	A G	G C	A G	T G	T G	G G	A G	G C	T T	A G	T G	G C	T C	G A	T G	G G

FIG. 5a.

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SEQ: SEQ: SEQ:	. ID. . ID.	NO NO	. 19 . 20	C G	C G	T G G T	G C	G C	G A	G G	G G	A C	G C	C T	G G	C	G C	A A	G	G C	T C	G C	G G	T C	G	G	G T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C G	C A	T C G T	A A	A T	G G	G G	T C	C G	C	G T	C G	A G	A A	G G	T G	G A	C C	C G	T T	G G	G A	C A	C T	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A G	C C	C G C A	G G	C C	T A	C G	C G	T G	G A	G C	A A	C T	A C	G C	A T	T G	A C	T C	G G	G G	A A	C	A T	C A
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A T	C G	T C A T	C G	A C	G T	C	C A	G A	C G	T C	G T	T C	G A	T T	C C	C	G A	A C	A C	T A	C C	T G	G A	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	C G	C C C G	A A	A A	G G	T T	C G	T T	T G	A A	T T	T C	T C	G A	A G	C G	C C	C C	T A	G A	G G	A C	A C	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A C	T C	G G A	G A	G G	A T	A A	G C	G C	T T	T A	T T	T A	C T	C G	T A	G G	A C	C T	G G	G C	G T	T C	G T	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G C	G A	A A	C C	C G	T A	C C	C C	C C	A T	G A	C T	T C	C A	T A	G G	G A	A T	C C	G A	G	A C	G C	C T	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C A	G T	G G	G C	T C	G T	G G	A G	T C	T T	T G	C C	C A	C G	G C	T T	G C	T	G G	A T	C	C T	C C	C C	G A

FIG. 5b.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A C	C C G C	T C	T T	C G	C G	A T	T G	C G	T C	G T	G G	T A	G G	G G	G C	C T	A G	G C	C T	T A	C G	C G	C A	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G G	G A T G	G G	C G	A A	T A	C C	T C	G T	T C	A A	G T	T T	C G	A T	G G	G C	G T	C T	C T	A C	G C	T T	G A	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A G	C G G T	C C	A T	C C	C C	C A	C G	C	A T	A C	G A	C C	C C	C A	C G	A C	C C	T C	G T	C G	C T	A C	G A	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	C G C C	A C	A G	T G	C C	G A	A G	A C	C G	G T	C T	C T	A C	C C	A C	C C	T A	C C	A T	G T	A T	A C	C T	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G C	G C C A	G G	C A	G A	C C	A G	G C	T A	G C	T C	A C	C A	A T	T C	C A	G G	G C	G C	G A	C C	A A	C	T T	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T C	A T A T	T C	C A	C A	C C	A C	T C	G T	A A	G C	C C	C	G G	G C	G G	G T	C G	T A	G A	G A	C C	C T	A C	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G T	T	G G	G A	C A	C A	A A	G G	G T	C G	C G	T G	G G	C	C T	A G	G G	C A	C A	C G	G A	C A	G G	G A	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G T	G G	A C	G T	A A	T C	G C	G A	C T	G C	C	T A	G G	G C	A A	G G	G A	A C	C C	G A	T C	G T	A G	A A	T G

FIG. 5c.

SEQ. SEQ. SEQ.	ID.	NO.	19 20	A G	G T	C	C T	G T	C C	A A	G C	G T	G T	A C	C	A A	T C	C T	C C	T T	G G	.C G	C A	G	G	A A	GCCC	T
SEQ. SEQ. SEQ. SEQ.	ID.	NO. NO.	19 20	A T	T G	G G	A A	G G	C G	T A	C A	A C	A G	G A	C	T T	C G	A A	T A	C G	C G	A A	C G	C G	A C	C T	0000	A G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C A	A A	G T	C T	A G	A A	G G	T A	G T	T T	G A	A C	T T	C T	C T	A C	G C	G G	C C	C	A A	A G	G A	A C G A	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A T	C T	C C	A T	A T	G C	T T	A C	C A	C G	T A	A T	T C	A C	T A	G G	A C	G T	C G	T T	G G	C C	T C	G C C C	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A T	C C	A A	A A	C A	G A	A A	C C	C C	C T	T G	A A	T A	C G	A C	A G	G C	A C	T A	C G	A G	T A	C T	C C G A	T C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T C	A C	T G	G A	C A	C T	T C	G A	G T	C C	T G	G T	C G	A G	G G	C A	T C	C T	T T	G	T T	C C	T T	A C A T	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A G	C A	G G	C A	T C	G T	G G	T A	G A	G G	C	T C	G C	A G	G G	G A	C A	T A	G G	C T	T T	A T	G T	G T	A T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T G	G T	T G	G A	G G	A G	A T	C G	C T	T A	C	A A	T A	T G	G	T A	G G	C C	T G	T T	T C	C T	C C	T T	A T

FIG. 5d.

SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T T	G G	G G	C G	T A	C A	C G	A A	G A	C G	T T	C A	A C	C G	C T	A C	G T	C G	C G	C T	T T	G C	T C	C T	T A C T
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A A	A T	C T	G C G T	G G	G G	C T	A G	G G	C T	G A	T T	T G	T C	C T	C	C A	C	A A	C A	T T	T T	T G	C G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T T	C C	C A	G G A A	A G	A A	C T	G C	C T	A A	C C	C G	C A	A C	T C	C C	A T	G T	C C	C T	A A	C T	A C	C A	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C C	C T	A G	0000	A A	A C	C A	C G	C T	T G	A G	C A	C T	C G	G A	C G	G A	T T	G	A A	A C	A T	C	T A	C G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T G	T C	T G	T G G T	A T	A G	A G	A A	G G	T G	G G	G C	G C	G A	C C	T A	G T	G C	A A	A C	G A	A A	A C	G T	A G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	T G	G A	A C T C	T T	A G	C T	C C	A A	T T	C G	C C	A T	G G	C A	A A	G T	A C	C C	C T	A G	C C	T C	G A	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G T	G A	T C	C C	T C	T G	C C	A A	C G	T C	T A	C T	G T	A T	C C	T C	C A	T A	G C	G A	A T	C G	G A	A C	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C T	T C	G C	G C	A A	G G	G G	A A	A A	C T	G T	A T	G G	T T	G G	A G	A A	G G	G A	A A	G A	G C	C T	T A	G A

FIG. 5e.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G C	G A C T	A A	T A	T G	G C	A G	G A	A C	T T	T	A A	C A	T A	T A	T G	C A	C	G A	0	C	A C	G T	A G	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T G	G T G A	T A	C G	T A	T C	C A	T G	C G	A A	G G	A G	T C	C T	C T	A C	G C	C A	T G	G G	T A	G G	C G	C C	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G C	T C C	C G	A C	A T	A G	A G	A C	C C	C T	T A	G T	A G	A A	G T	C	G C	C	C A	A T	G C	G T	A G	T G	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C C	0 0 0	C T	G T	A G	A G	T C	C A	A C	T T	C G	G G	T C	G C	G C	G T	A G	C A	T A	T C	T A	T A	C G	T A	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	T G T A	A C	G T	A G	C G	T A	G G	A G	A A	G G	C G	C C	C G	G G	G C	A C	A G	A T	G T	T C	T T	T G	T G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T G	G G T T	T G	G C	A G	G C	G C	T T	G G	T G	A A	C G	A G	A A	G C	G T	A T	G C	C A	G A	T C	C T	T A	C	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	С	G A	G A	G C	A C	A A	G G	A A	A C	G C	T A	A T	C T	G A	T C	C C	T G	G A	G C	T C	T A	C A	C A	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID	NO.	19	С	A T	T A	T	G C	G	G G	T	G	G A	T A	A T	T	G A	C A	T	G T	A C	C T	A T	A C	T	T T	G C	C C

FIG. 5f.

SEQ SEQ SEQ SEQ	. ID . ID	. NO . NO	. 19 . 20	T T	T T	C T	A G	A A	G G	A G	T G	C T	T	A T	C	G	C	C T	C G	C G	T	T	C A	T T	A G	T T	C	A A G T
SEQ. SEQ. SEQ.	. ID.	NO. NO.	19 20	A T	C G	T T	G T	C T	A G	C A	A T	G G	T C	C	G A	A G	T C	G G	A G	G C	A T	T C	G T	A C	C	T	G A	C A T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	19 20	G G	G	C C	G A	G T	T G	G G	G A	A C	G	G C	G T	C T	C A	A T	C C	A G	T A	C G	A C	C A	A G	A C	A C T C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G C	A A	G G	A G	T G	T T	G G	T G	C C	A A	T G	C	L C	T A	G C	Α Λ	A A	T G	С Л	C A	T G	G A	C T	C T	
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A G	T C	A T	C A	C C	C T	G A	C T	A G	G A	C	A A	T G	T C	T A	C C	C C	A A	A A	C G	A G	T A	G T	G A G T	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A T	T C	C T	C T	C T	A C	G C	G T	A G	A G	T T	T C	T C	G A	T A	G A	G A	A C	G A	A G	A A	A T	C A	C T A	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	19 20	A T	C	C G	A A	A T	G T	C G	G G	A A	C G	T G	G G	A T	A C	A C	A C	G C	A C	C	A C	C A	C G	C	G T T C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A A	G C	G C	A A	G G	A A	C	A C	G C	G T	A G	G	G T	C C	T A	T T	C	C A	A A	G G	G A	A C	G A	C G T A	C T

FIG. 5g.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	19 20	A C	C C	C G	0 0 0 0	C T	T T	G C	G C	C T	C	T T	A C	T A	G C	A A	T G	G A	C A	C A	A C	T T	C	T T	G T	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G A	C T	C	G T T A	T C	G C	G G	C T	A C	C T	T C	G A	G G	C T	C T	C C	T T	G C	A T	A C	C C	A A	A G	G C	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C T	A G	T G	T C G T	T C	G A	G T	A T	G G	G T	A C	G C	G Ţ	C A	G G	G C	C T	C G	G T	T T	T G	C T	T C	G T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T T	G C	T T	C G G	C T	G C	C C	C T	T T	G T	G A	A A	G C	G A	A T	C C	T T	T A	C C	A A	A A	C C	T T	A C	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A C	A A	C T	T A G T	A T	C	C C	A G	G T	A T	C A	C T	A A	T T	T C	A C	C A	C G	G A	A A	C C	C T	A C	A A	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	C G	T C	A A C A	C C	C A	G A	G C	G C	C T	A G	A A	T A	G C	A A	A A	C	T C	C T	T G	T A	C C	G T	T G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C T	T G	T T	G	G G	A G	G C	G T	G G	T C	G T	T C	C A	T C	C T	T G	G G	G C	C T	C T	A T	T A	G	T C	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G	T C	G T	T G	T T	T C	G T	A T	T C	G C	C	C C	A C	G T	C G	G G	G G	C G	T C	C T	T C	C G	G A	G T	A G

FIG. 5h.

SEQ. II SEQ. II SEQ. II	D. NC D. NC). 19). 20	T G	G T	G T	C A	A C	T C	G A	G C	A A	T	G T	C G	T G	T G	A A	T G	C G	G A	A A	C C	C	A A	C G	T C T T	T T
SEQ. II SEQ. II SEQ. II	D. NC D. NC). 19). 20	T T	C	A C	G T	G T	G T	T C	G G	G T	C C	A T	G G	C C	T C	A A	C G	A G	A C	G C	A C	A G	G C	A C	C T T A	T C
SEQ. IL SEQ. IL SEQ. IL	D. NC D. NC). 19). 20	G T	G G	C G	T C	A T	C C	T C	A T	T G	G G	A G	C C	A C	G T	C G	A G	C G	C C	A T	A T	G T	G A	A G	G T T G	G C
SEQ. II SEQ. II SEQ. II SEQ. II	D. NC D. NC). 19). 20	A T	T	C C	T G	T C	T T	C A	C C	T G	G G	G T	T T	C	C	A A	A T	A G	A T	C T	A C	G A	A C	T C	G A A G	A A
SEQ. 11 SEQ. 11 SEQ. 11 SEQ. 11	D. NC D. NC). 19). 20	A G	T A	G T	G T	A T	T G	T G	G T	G G	A G	G G	G T	G C	T C	C A	C C	C A	C C	C G	C G	C T	A C	G T	A C T C	T C
SEQ. II SEQ. II SEQ. II SEQ. II	D. NC D. NC). 19). 20	G A	A C	C A	C A	A A	G G	A A	C A	C G	C G	T A	G A	G G	T A	C A	A A	T A	C	A A	A A	G G	A G	C A	C A G A	T T
SEQ. II SEQ. II SEQ. II SEQ. II	D. NC D. NC). 19). 20	T G	C G	C A	G G	C G	T A	T A	C G	C A	T C	G T	T C	C T	A G	G C	A A	G A	A C	A C	A C	C T	G G	C G	T A	T A
SEQ. II SEQ. II SEQ. II	D. NC D. NC). 19). 20	T G	A C	T T	C	T T	C A	C T	GGC	T C	C C G	7 A	C C	A A	G	T T	T G	C	T G	C C	T C	C T	C G	A C	G T	C G

SEQ. ID SEQ. ID SEQ. ID SEQ. ID	. NO. . NO.	19 20	C	T T	G	G G	A G G G	C	A A	T T	T G	G G	T A	C T	C	T	A C	G C	C T	T	G A	T	T	C C	T T	C	Ţ
SEQ. ID. SEQ. ID. SEQ. ID. SEQ. ID.	NO.	19 20	G C	T C	C A	T T	G G C T	T T	C G	C G	T C	T A	T G	A A	A T	C C	A G	T T	C G	T G	A A	C	A C	A C	C T	T C	C T
SEQ. ID. SEQ. ID. SEQ. ID. SEQ. ID.	NO. NO.	19 20	A G	C C	A A	T C	T G C G	T G	C G	C A	G C	T C	T A	A T	T T	A G	Т А	C G	C A	A C	G A	A T	A T	C T	T G	C	A C
SEQ. ID. SEQ. ID. SEQ. ID. SEQ. ID.	NO. NO.	19 20	Α	A A	G G	C G	C C A A	C G	A G	A A	C A	C C	T C	G T	A A	A A	C G	A G	A A	C A	C G	T A	G T	A A	C T	T T	G G
SEQ. ID. SEQ. ID. SEQ. ID. SEQ. ID.	NO.	19 20	Α	T C	G G	T T	G	G T	G C	C T	T A	G T	C T	T C	C T	A G	C	T C	G C	G C	C A	T G	T C	T T	A G	G G	C A
SEQ. ID. SEQ. ID. SEQ. ID. SEQ. ID.	NO.	19 20	G	G C	C A	T T	A G T C	T G	C	T A	T G	C	C T	C C	C C	C A	T G	G G	G A	G A	G	C A	T T	C G	G A	A A	T T
SEQ. ID. SEQ. ID. SEQ. ID. SEQ. ID.	NO.	19 20	G A	G C	T A	T T	A G	C G	C	A T	C T	G G	T G	T C	G A	G T	G	A T	G T	C C	A T	A A	C T	C	A G	G T	T T
SEQ. ID. SEQ. ID. SEQ. ID. SEQ. ID.	NC. NO.	19 20	T A	T C	C A	C A	T G	T G	T G	C G	G C	T T	C G	T C	G T	C G	C	A T	G G	G C	C T	C G	C	G T	C G	C G	T G

FIG. 5j.

SEQ SEQ SEQ	. ID . ID	. N O . N O	. 19 . 20	C A	T A	G T	G C	C T	T T	C	C	T T	G	G G	G C	C T	C	T A	G T	G G	G A	C G	T A	T	T C	A A	Α	T
SEQ. SEQ. SEQ. SEQ.	. ID. . ID.	NO.	. 19 . 20	C A	T G	G T	G G	G T	C G	T T	A C	C C	G A	G C	T T	T G	C A	C G	A A	T A	G G	T A	T T	C	A A	C A	C C T C	A G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A A	G T	A C	T A	T C	T C	G G	G G	T G	G C	G T	G G	T T	C G	C G	A G	C C	A A	C T	G G	G G	T C	C T	А Т А А	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C C	A T	C A	A C	A A	A A	G T	A G	A T	G G	G G	A C	A A	G G	A T	A C	A C	A T	G G	A T	A G	G C	G C	A A T G	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	G T	G C	A A	G C	G T	A G	A C	G T	A C	C C	T T	C G	T T	G C	G A	A C	A C	C A	C T	C G	T A	G T	C G T	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A T	G G	C T	T C	G C	T A	A G	T C	G C	C A	C G	A C	C A	A G	G G	T A	G T	G G	G C	C A	C G	T C	G C	A C T C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G T	G G	T C	G C	G T	G T	C T	A G	T C	G C	G T	A C	T T	G C	T T	C T	C G	T C	C C	A A	C T	T A	C G	T T	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G T	C T	C C	A T	T C	C C	T T	G C	G C	C T	A A	G T	A A	T T	C C	G A	T C	G T	G C	A T	C T	C G	C T	T i	C G

FIG. 5k.

T G G C A G C C	G C T T A G	G G A G C	Α .	T G	G A	A T	T
T (A .	A C	C C G A C C	T T G C G A A C	T C T C A C	CA	A A ^ a
T	T G	C A	T A	T +	C '	C (
T C	A G	G G	A A	A T	G A	A	T
A G	G T	A T	G C	T T	T A	G G	A
C T	A A	C A	A A	C G	C G	A A	G A
A A	A A	C	A A	T	G	G G	A A
G G	G G	C C	C C	T C	T T	T G	A C
A A	G G	C A	G A	T G	C C	A A	G T
G A	A G	G C	A A	T G	G A	T G	A A
T C	A G	T A	C	A C	T A	T A	C C
T C	T A	C G	C C	C C	C G	C A	T C
A C	C G	T G	T A	G C	G T	G A	C G A
C G	C	T A	C	G T	G	T G	A C C
C	A C	A C	C C	T	C	T A	C G C
A G	A C	T G	A T	T A	G C	C G	C T A
G T	G A	C	C A	C A	G A	C T	T C A
G T	G C	T G	C	G G	G A	T C	G A A
C T	A T	C G	T T	G A	G G	G	T A
C	G A	T A	T G	G	A A	T	G G
A T	G G	G G	G A	A G	A G	T	T T
C	A T	C G	C G	C A	C G	A A	G C
G	A C	A C	G A	A G	A A	A C	A T
T	C G	G T	A C	T G	T A	T	C
. 19 . 20	19 20	19 20	19 20	19 20	19 20	18 19 20 21	18 19 20 21
NO NO NO NO	NO. NO. NO. NO.	NO. NO. NO. NO.	NO. NO. NO. NO.	NO. NO. NO. NO.	NO. NO. NO. NO.	NO.	VO. VO.
ID. ID.	ID. ID.	ID. ID.	ID. ID.	ID. ID.	ID. ID.	ID. ID	ID. (ID. (
SEQ	SEQ. SEQ.	SEQ. SEQ.	EQ. EQ.	EQ. EQ.	EQ. EQ.	EQ.	EQ.
	5	9	5	5	S	S	S

SEQ. SEQ. SEQ.	. ID. . ID.	NO.	19 20	G T	A C	C T G	C	A G	C G	C	G A	G G	G C	C A	T G	G C	T T	G C	G C	G G	C C	A T	T C	G C	G C	C G	T G	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NC.	19 20	T G	C C	A T C G	A A	C C	A C	A C	T A	G C	T C	G G	G A	C	A A	G C	T C	C C	C C	T C	G A	T G	G A	C A	C C	T C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C	A T	G T C A	C T	A G	C G	T G	G G	C G	T C	C C	C T	T G	G C	T C	C C	A A	C G	C G	A G	T G	G A	A C	T C	T C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C C	T C	C G T T	T G	C A	C G	A C	G C	C C	C	A C	G C	C G	A A	G C	G C	A G	T G	G C	C T	A T	G A	C G	C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T G	T T	G G G A	C A	C T	T	T G	T G	G A	C G	C T	T C	C G	T A	C G	T T	T G	G C	C A	C T	A T	T T	A G	G C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T T	T T	C T A T	C T	T A	C A	C G	T T	C G	C A	T G	A G	T G	A T	T A	C G	A G	C G	T T	C G	T A	T G	G G	T G	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G G	T G	C G A G	C C	T A	C G	T G	T C	T C	G A	T G	G T	C A	C G	C G	A G	A G	G G	A A	T G	G G	C G	G A	C A	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G	G G	A C G C	T A	G G	A A	T G	C G	A G	C G	C A	C A	G G	A G	G G	G C	G A	G G	A G	A G	T G	G A	C C	C T	A C

FIG. 5m.

SEQ. SEQ. SEQ. SEQ.	ID.	NO. NO.	19 20	G A	T G	C G	G A	G A	A G	G C	G A	C G	G G	C G	A G	G G	G T	A C	C	A C	C C	T C A A	A T	T C	C	A C	A C	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A G	C C	A T	G G	G G	G G	T A	C A	A G	T A	C A	G C	A A	C T	C G	A C	A T	C A	A A T A	A C	C	A A	A A	C T	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A T	G C	G A	A T	G C	G T	A C	G T	A T	A G	G T	T A	C A	C A	C T	G A	C	C A	G T G	G	T T	T C	G C	G C	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C C	A T	A G	G T	G G	A A	G G	A T	A T	C	C T	C G	T G	C	A C	A T	C G	T A	G G T C	G T	A T	A G	A G	A G	G T
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	A C	T T	C C	A T	T C	T A	G T	C A	T C	G C	A T	G C	A T	A G	A G	G G	A A	G A	G G A A	A C	G A	C G	G A	T C	C G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T T	C T	T T	C T	T T	G C	A T	A C	C T	T C	G T	C T	G A	C C	C T	A G	T C	C T	G A T T	G C	C A	T T	C	C T	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	G A	T T	C T	T T	C T	G G	G T	C A	A T	G C	C A	A C	C C	C T	T C	C T	C T	G C	Α	T C	C A	C A	C T	G T	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C A	G	C T	C T	A C	C G	C T	C A	A C	C	C T	G G	A G	C C	A T	C T	C	C A	C A	C G	A C	G T	A G	A C	C T

FIG. 5n.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C C	C A	T C	C T	T G	G C	G T	G C	G A	G C	C A	C C	T G	G C	C T	C G	C	A C	G T	C	G C	G T	A C	G C A T	C G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	C	C A	C G	T C	G C	A T	G C	C A	C C	C T	C G	C C	C A	G T	A C	C T	C T	G T	G C	C T	T	T T	A T	T G C G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T C	G A	T T	G G	A C	T A	G A	G C	G A	A C	G C	T C	C T	G C	A T	G T	T C	G T	C A	A G	T T	Ţ Ţ	T A	G G C U	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	T A	T C	T G	A G	T C	A A	A A	G C	T C	G C	A C	G T	G	G	T	A	G	G	G	Т	G	Α	G	A G A	G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	Α	G	G	Α	С	A	G	G	С	С	Α	G	T	A	G	G	G	G	G	Α	G	G	G	T A C	Α
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	Α	G	G	G	Α	G	A	G	G	G	G	Α	Α	G	G	G	С	Α	G	G	G	G	Α	G C	T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	19 20	С	Α	G	G	Α	Α	G	С	Α	G	G	G	G	G	T	С	С	С	С	Α	Τ	С	С		С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	Α	G	С	T	G	G	G G	A G	Α	G C	A A	A G	C G	Α	T	G	С	T	Α	T	С	С	Α	A	T

FIG. 50.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	С	A T C	С	Α	Т	С	T	С	T	T	G	T	Α	A	Α	T	Α	С	Α	T	G	T	С	С	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	С	A C	T	G	T	G	A	G	T	T	С	T	G	G	G	С	T	G	Α	T	T	T	G	G	G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	Т	T C	T	С	T	С	Α	T	Α	С	С	T	С	T	G	G	G	Α	A	Α	С	Α	G	Α	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	С	C T	T	T	T	T	С	T	С	T	С	Т	Т	Α	С	I	G	С	T	T	С	A	T	G	Т
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	Α	A A G	T	Ţ	Т	Ţ	G	T	Α	T	С	Α	С	С	T	С	T	Т	С	Α	С	Α	Α	Т	T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20	Т	G A A	G	T	T	С	G	T	Α	С	С	T													
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20																									
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	19 20		A C																							

FIG. 5p.

SEQ. ID. NO. 18 AGGCAGCAGTTACAGAAACGTA SEQ. ID. NO. 19 CCATGCAACACCCTCTTCTAGTTAC SEQ. ID. NO. 20 SEQ. ID. NO. 21 SEQ. ID. NO. 18 GTGAATTCA SEQ. ID. NO. 19 CACGGCAACCCCTGCAGCTCCTG SEQ. ID. NO. 20 SEQ. ID. NO. 21 SEQ. ID. NO. 18 SEQ. ID. NO. 19 CCTTTGTGCTCTGTTCCTGTCCAGC SEQ. ID. NO. 20 SEQ. ID. NO. 21 SEQ. ID. NO. 18 SEQ. ID. NO. 19 AGGGGTCTCCCAACAAGTGCTCTTT SEQ. ID. NO. 20 SEQ. ID. NO. 21 SEQ. ID. NO. 18 SEQ. ID. NO. 19 CCACCCCAAAGGGGCCTCTCCTTT SEQ. ID. NO. 20 SEQ. ID. NO. 21 SEQ. ID. NO. 18 SEQ. ID. NO. 19 CTCCACTGTCATAATCTCTTTCCAT SEQ. ID. NO. 20 SEQ. ID. NO. 21 SEQ. ID. NO. 18 SEQ. ID. NO. 19 CTTACTTGCCCTTCTATACTTCTC SEQ. ID. NO. 20 SEQ. ID. NO. 21 SEQ. ID. NO. 18 SEQ. ID. NO. 19 A C A T G T G G C T C C C C T G A A T T T T G C SEQ. ID. NO. 20 SEQ. ID. NO. 21

FIG. 5q.

```
SEQ. ID. NO. 18
SEQ. ID. NO. 20
SEQ. ID. NO. 21

SEQ. ID. NO. 18
SEQ. ID. NO. 19
C C A A G G T C A C A T G C T C C C T T G C C T C SEQ. ID. NO. 21

SEQ. ID. NO. 19
SEQ. ID. NO. 20
SEQ. ID. NO. 18
SEQ. ID. NO. 18
SEQ. ID. NO. 19
SEQ. ID. NO. 19
SEQ. ID. NO. 19
SEQ. ID. NO. 20
SEQ. ID. NO. 21
```

FIG. 5r.

SEO, ID. NO. 23 A T G G G C C C G G G G G G A C C C T G T A C C C C A G T G SEO. ID. NO. 22 CTCCGCCCCCTGGGCGCTGGCGGGGGCAG SEQ. ID. NO. 23 GGGTGGCCGCTGCCTCTTCTGCTGGTGATG SEQ. ID. NO. 22 ACCCCAACGCCACCTCGGAAGGTTGCCAG SEQ. ID. NO. 23 GCGGCTGGGGTGGCTCCGGTGTGGGCCTCT SEQ. ID. NO. 22 ATTATACATCCGCCCTGGGAAGGTGGCATC SEQ. ID. NO. 23 CACTCCCCTCATCTCCCGGGGCCTCACCCG SEQ. ID. NO. 22 A G G T A C C G T G G C T T G A C T C G C G A C C A G G T G SEQ. ID. NO. 23 AGGGTCCCCCCGCACCCCTCCTCAGAACGG SEQ. ID. NO. 22 A A G G C C A T C A A C T T C C T G C C T G T G G A C T A T SEQ. ID. NO. 23 CGTGCAGTATACATCGGGGCGCTGTTTCCC SEO. ID. NO. 22 GAGATCGAATATGTGTGCCGAGGGGAGCGC SEO. ID. NO. 23 A T G A G C G G G G C T G G C C G G G G G C C A G G C C SEQ. ID. NO. 22 GAGGTGGTGGGCCCAAGGTGCGCAAATGC SEQ. ID. NO. 23 TGCCAGCCCGGGGGGAGATGGCCGCTGGAG SEQ. ID. NO. 22 CTGGCCAACGGCTCCTGGACGGATATGGAC SEQ. ID. NO. 23 GACGTTAACAGCCGCAGAGACATCCTGCCG SEQ. ID. NO. 22 A C A C C C A G C C G C T G T G T C C G A A T C T G C T C C SEQ. ID. NO. 23 GACTACGAGCTCAAGCTTATCCACCACGAC SEQ. ID. NO. 22 AAGTCTTATTTGACCCTGGAAAATGGGAAG SEQ. ID. NO. 23 AGCAAGTGTGACCCAGGGCAAGCCAAAG SEQ. ID. NO. 22 GTTTTCCTGACGGGTGGGGACCTCCCAGCT SEQ. ID. NO. 23 TACTTGTACGAACTACTCTACAATGACCCC

FIG. 6a.

SEC SEC	. ID . ID	. NC . NC). 22). 23	A	T	G	A	A A	T G	G A	G	A C	Ā	CT	C	C	G	C	G A	7	G	C	A C	G -	T	- G	C	C	G	Ā	τ Α	G	7
SEC SEC																																	
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SEQ. SEQ.	ID. ID.	NO. NO.	22 23	C G	G T	CC	A T	G T	A C	G A	A C	C	A T	T C	C A	C	T	G G	C	C T	G	G	A A	C	T G	A A	C	G C	A T	G G	C G	T A	C G
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A G	A A	G G	C	T G	T A	A G	T T	C G	C A	A A	C A	C G	A A	C G	G G	A C	C T	A G	G G	C G	A A	A T	G C	T G	G A	T	G A	A T	C
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	C A	0	A T	G T	G T	G C	C	A G	A A	G C	C A	C G	A A	G	C T	A T	A T	G	Ţ	A T	C	T T	T	G	T	A A	C	G C	A C	A A
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	C G	T	A T	C	T T	C G	T C	A C	C	A G	A T	T T	G A	A A	C A	C A	C	0	A C	T T	C G	A A	A A	G G	A C	T G	C	A C	T A	Ŧ A
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	C	T A	C	A G	T C	G T	C	G	A	A	Т	С	т А	T	С	A G	G T	T	T G	C G	T A	G C	<u>T</u>	C T	T T	C	CC	A T	C	^

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SEQ. SEQ.	ID. ID.	NO. NO.	<u>22</u> 23	C	T A	T G	G A	T	A G	G G	C A	T A	G	A C	GC	G	CG	T	G A	C	C A	C	G	G T	<u>^</u>	T	G	T	G	G	A G	A A	C G
SEQ. SEQ.	ID. ID.	NO. NO.	. 22	C	T	TC	A T	T A	T	G A	T A	G G	C G	T A	C A	T A	C	A G	7 C	A T	T C	G T	G	C	J- G	C	C	A A	G A	T G	T A	C A	A G
SEQ. SEQ.	10. 10.	NO. NO.	. 22	C T	C	A C	G G	C	00	T T	T G	G G	T	C T	A C	A C	A T	CC	C A	G T	A C	C	A G	G G	C	G G	G	T	T A	T T	C	CC	C T
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A G	C A	G C	T A	T A	СС	T	T G	C G	C T	G	G C	A A	C A	G G	C A	A C	T C	C	C A	A T	T G	C A	С	G C	СС	C G	A T	C C	A A
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	C A	T	СС	C A	A A	C	A T	A G	T	C A	CC	C A	A G	C T	C G	CG	G A	G A	G	T A	G A	A A	A T	A G	C A	T C	C	T G	T A	C G
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G G	A C	A G	A G	A T	G G	T G	G A	G G	G G	G G	0 0	T C	G A	G C	A A	A T	G C	A A	A C	G C	A A	T C	C G	G G	C A	T G	A A	C T	C T
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A G	T	СС	C A	A T	A G	CC	A T	G G	A A	C A	СС	A C	CC	C	G G	A C	G C	G A	T A	CC	T A	T	CC	A C	C	C A	T A	C G	A C
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A A	C	G	C	T	G	G A	A A	T C	G A	A T	C G	C A	7 C	G G	G T	A C	G A	G C	A A	G G	C	G A	A	G T	T	G T	A G	A T	A G
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G G	A A	G	G A	C A	T A	G C	G T	G A	A A	T	C	G A	A A	G	A C	T G	C G	A C	C T	T	T A	T A	C A	C A	G G	A A	CC	A A	G C
SEQ. SEQ.	ID. I D.	NO. NO.	22 23	A C	G C	T	T G	T A	C	T G	T A	C G	T A	C	G T	G G	A G	T A	C G	G	A C	G T	C T	T	G C	T A	G	C G	C A	T	G	T	T A
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A C	A C	A A	A C	A T	C G	C G	T C	G	A T	A A	G	C G	G A	T	C	A C	A T	G A	A T	T C	G	C	T	C	GC	A C	A T	T T	C
SEQ. SEQ.																																	
SEQ. SEQ.												T	T	С	С		G	С	G						TC								

SEQ SEQ	. ID. . ID.	NC. NO.	. 22 . 23	A	G T	G	C A	T A	00	- - -	Ŧ	70	G A	G A	G	A	A A	G	A C	A	G	T A	A C	CC	G	T T	C + ·	Ŧ	G	G	G	Ŧ A	СС
SEQ SEQ																																	
SEQ.	. ID. . ID.	NO. NO.	. 22 . 23	A T	A T	G	A G	C A	CG	T	A G	T	G	A T	C	C T	C	G	T	C	A C	A C	T A	C	A G	A T	G	T	G T	T	A T	C	A T
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SEQ. SEQ.																																	
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A C	A A	CC	C	A T	G T	A C	C G	T	T _	Α	エエ(フ	Α	T	С	С	A A	C G	C A	A A	G C	A T	T	00	- C	A	C G	CC	G C	G C

SEO. ID SEO. ID). NO). NO	. 22 . 23	G A	۵		C (A C	T T	G G	A A	A A	C	T A	C A	C	T C	C T	C	T A	СС	CT	T G	T	T	G G	A T	G	G G	G G	C	G	T	T C
SEQ. IE SEQ. IE					; ;	T (G	G	C	C G	A C	T A	G C	T T	G G	G G	T C	C	T G	T	-	G G	A T	T C	G T	C	CC	A C	G C	C	G C	G T	C
SEQ. IL SEQ. IL				· (C (C G	C C	G T	G G	A G	T A	G T	G	C G	A T	T	G A	G C	A C	C A	A C	C A	T	T A	A G	T G	C G	G A	A G	G A	C A	A G	G C
SEQ. IL SEQ. IL	D. NC D. NC). 22). 23		;	Γ,	A G	C T	A T	G	G C	G C	C G	G T	G T	C	A G	G T	СС	T	A G	C C	A C	A A	G G	A G	A C	G C	A C	T G	СС	G	G T	C
SEQ. IL SEQ. IL	D. NC D. NC). <i>22</i>). <i>23</i>	ר י ד		4 (3 (C G	T C	A T	C	G	A T	C G	A G	G G	CC	A T	C	C G	A G	A G	G C	G T	A T	T	G A	A G	T	СС	T	T G	T G	C	СС
SEQ. II SEQ. II				· /	3 (G T	T G	C	CC	A T	A C	A T	A A	C T	G G	G T	A T	CC	A	A C	G C	T A	G A	G G	A	7	T	G T	G	A G	G T	G	G G
SEQ. IL SEQ. IL	D. NC D. NC). 22). 23	? 7	- (C T	T C	C C	C	C	C A	CC	A A	G	C	T	G T	A T	CC	C A	A C	G G	A A	C A	C	T A	T A	G	G	T A	C	A G	T A	C G
SEQ. IL SEQ. IL	D. NC D. NC). 22). 23	? A		۹ <i>ا</i>	G G	A A	C	A G	T	T A	C G	C	G G	T	T A	T G	C G	C A	T A	G G	T A	СС	T C	CC	A T	G A	A G	A A	A G	СС	T C	C
SEQ. II SEQ. II				- (T G	T G	A A	T A	C A	T	C	CC	G	T A	C T	T	СС	A C	G A	7 C	T	C G	T	C G	T	C G	CC	A C	G	C	00	T	G
SEÒ. II SEQ. II			? (à (G T	C G	A G	T	T	G A	T	T G	C	T A	T	G G	C	T C	G	T	T G	G A	T	C T	T C	G	T	C	T	G	T A	C	C
SEQ. II SEQ. II	D. NO D. NO	o. 22 o. 23	? -	Γ :	T G	T G	A C	A	C	A A	T T	C	T G	A T	C G	A G	A A	C	T	C	CC	C	A T	C G	G	T A	T	C	G	T A	T A	A C	TC
SEQ. II SEQ. II	D. NO D. NO	D. 22 D. 23	2 ,	۹ ۹	T T	C T	C G	A A	G	A	A	C	T	C	C	C	A	G	A	. C	C	A G	A A	C	C	T A	G A	A C	A C	C A	: А . А	A	T G
SEQ. I. SEQ. I.	D. NO	0. 2: 0. 2:	2 (C G	T A	G A	A G	C A	T	G	. T	: T	G	ìΑ	T	G	T	C	7	. C	; C	; T	C	: A T	C	: T	G	i C	S C	; A		: T : A	G G

SEQ. SEQ.				G	C	T	G G	C A	T G	G C	Ŧ	CC	-	T G	C	C A	C G	T (C	C	5- C	C	G A	G A	G	C A	Ŧ	G G	G A	* * • • • • • • • • • • • • • • • • • • •	÷ G	G A	G A	
SEQ. SEQ.				T A	A C	C G	C T	A G	C G	A C	T	A T	G	G G	G C	A A	G	A	A T	G	CC	C	A A	G	T G	- G	C	C	C	G	T A	T A	T G
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G	T G	C	TC	G T	C	CC	A T	G	G C	C	C	00	G	C G	C	T T	T G	T G	G	G A	C	T	СС	-	- - - -	G	G C	G	CT
SEQ. SEQ.				T G	T	G	G T	G A	C	T	T A	T A	A A	G C	T	C A	T A	G G	G A	G	CC	T	A	T G	G T	G C	СС	- A	0 0	1	A G	T A	G A
SEO. SEO.	ID. ID.	NO. NO.	22 23	T A	T A	C	A A	C	CC	A A	A A	G T	A G	T A	CC	+ C	G A	G C	T A	G G	G G	G G	T	CC	CG	A T	CG	A G	C	AC	G A	T T	C
SEO. SEO.	ID. ID.	NO. NO.	22 23	T G	T	C	A A	C	G C	A T	A A	G	A A	A A	G T	G	A T	G C	G	A C	G G	A G	A T	G C	A C	A T	G G	G	A G	G	T C	G	C
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A A	G T	G	A A	A C	G T	A G	CC	C	CC	TC	A T	G G	A T	G	C A	C	C	T A	G T	G G	A A	A T	A	CC	T	C	T	A C	T C
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G A	C G	C	A C	C A	T	G	T A	G G	G	G	C	C G	TC	G A	C	T C	G C	G T	∓ ⊤	G T	G G	G C	C	A T	-	G T	G	A C	T
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G T	T C	C	0 0	T T	G	A G	CC	T	C A	T T	T	G G	C	C	A T	T T	СС	7	G	G T	C T	A C	GC	A T	Ŧ A	÷c	G A	T T	G C
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G A	A C	C	C	C	C	T G	T	G	CG	A T	C	CC	G	A C	A T	C T	C T	A G	T T	T	G	A C	C	A A	C A	T G	T A	T T	T G
SEQ. SEQ.																																	
SEQ. SEQ.																																	
SEQ. SEQ.											С	С	Α	A	С	Α	Α	С															
												F	-/	C	7.	6	5	/															

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SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A T	T	T C	T	T G	C A	T	A T	T G	G T	G	T	T G	A A	C G	A A	A A	G G	G	G A	G A	C A	Ŧ A	G	00	T	G A	C	T A	G A
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	CC	7 T	G G	C G	T A	G A	G A	G A	A G	A A	T	C	T A	T	T	C G	T C	T	G	C	G	Ā	<u> </u>	C	G	Â	A G	A G	C A	C
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	A C	A G	G C	A G	G T	СС	G T	T C	G T	T G	C A	C A	A C	C	T G	G	A G	A C	A C	A A	G T	A C	T A	C	A C	A T	T C	G C	A A	C G
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	C	A C	C	A C	G	G G	G C	C A	C	G	T A	G A	G C	G T	CC	A C	T G	G C	G	CC	T A	A C	T G	C	T C	A G	CC	A C	A A	T
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G C	TC	CC	GC	C	G A	G A	T	C A	CC	TC	C	T	G C	T A	C G	T A	C T	A C	TC	CC	A T	0 0	T	G	C G	T	C	C G	TC
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G C	T	G	A C	C	СС	A A	T	G	A G	T	C A	C	T C	T C	T	C	C T	A G	G A	T	00	A C	G	CC	A C	G	G G	A A	СС
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G C	C G	A G	G	C T	C T	T A	T	T C	G T	C G	C	T G	T A	T	G	C	C	T A	C	T	CC	T	G A	G	C	C A	A C	T A	C
SEQ. SEQ.	ID. ID.	NO. NO.	22 23	G T	T	G G	T C	T T	C	T T	C A	T C	T A	C A	C G	Т	Α	С	Α	Т	c	Α	С	Τ	С	T	G	G	Т	Т	G	T	G
SEQ. SEQ.				С	Т	С	7	Т	ī	G	T	G	С	С	С	Α	Α	G	Α	Т	G	С	G	С	Α	G	G.	С	Ŧ	G	A	Т	С
SEQ. SEQ.				Α	С	С	С	G	Α	G	G	G	G	A	A	T	G	G	С	Α	G	T	С	T	G	A	A	Α	С	G	С	A	G
SEQ. SEQ.				· G	Α	С	Α	С	С	A	Т	G	Α	A	A	A	С	Α	G	G	Α	T	С	A	T	С	С	Α	С	С	Α	Α	С
SEQ. SEQ.				A	A	С	Α	Α	С	G	Α	G	G	Α	Α	G	Α	G	Α	Α	G	Т	С	С	С	G	A	С	Т	G	ī	Т	G
SEQ.				G	Α	G	A	А	G	G	Α		^ 	_						A	Α	С	Т	G	G	A	Α	A	Α	G	A	T	С

FIG. 6h.

SEQ. ID. NO. 24 MILLLLVPLFLRPLGAGGAQTPNATSEGCQ SEQ. ID. NO. 25 MGPGGPCTPVGWPLPLLLVMAAGVAPVWAS SEQ. 1D. NO. 24 I I HPPWEGGIRYRGLTRDQVKAINFLPVDY SEQ. ID. NO. 25 HSPHLPRPHPRVPPHPSSERRAVYIGALFP SEQ. ID. NO. 24 E I E Y V C R G E R E V V G P K V R K C L A N G S W T D M D SEQ. ID. NO. 25 MSGGWPGGQACQPAVEMALEDVNSRRDILP SEQ. 1D. NO. 24 TPSRCVRICSKSYLTLENGKVFLTGGDLPA SEQ. 1D. NO. 25 DYELKLIHHDSKCDPGQATKYLYELLYNDP SEQ. ID. NO. 24 LDGARVEFRCDPDFHLVGSSRSVCSQGQWS SEQ. ID. NO. 25 IKIILMPGCSSVSTLVAEAARMWNLIVLSY SEQ. ID. NO. 24 TPKPHCQVNRTPHSERRAVYIGAL FPMS G.G. SEQ. 1D. NO. 25 GSSSPALSNRQRFPTFFRTHPSATLHNPTR SEQ. ID. NO. 24 WPGGQACQPAVEMALEDVNSRRDILPDYEL SEQ. ID. NO. 25 VKLFEKWGWKKIATIQQTTEVFTSTLDDLE SEQ. ID. NO. 24 KLIHHDSKCDPGQATKYLYELLYNDPIKII SEQ. ID. NO. 25 ERVKEAGIE ITFROSFFSDPAVPVKNLKRO SEO. ID. NO. 24 LMPGCSSVSTLVAEAARMWNLIVLSYGSSS SEQ. ID. NO. 25 DARIIVGLFYETEARKVFCEVYKERLFGKK SEQ. ID. NO. 24 PALS NRORFPTFFRTHPS ATLHNPTRVKLF SEQ. ID. NO. 25 YVWFLIGWYADNWFKTYDPSINCTVEEMTE SEQ. ID. NO. 24 EKWGWKKIATIQQTTEVFTSTLDDLEERVK SEQ. ID. NO. 25 A V E G H I T T E I V M L N P A N T R S ! S N M T S Q E F V SEQ. ID. NO. 24 EAGIEITFRQSFFSDPAVPVKNLKRQDAR! SEQ. ID. NO. 25 EKLTKRLKRHPEETGGFQEAPLAYDAIWAL

FIG. 7a.

SEC. SEC.	ם). ים.	NO. NO.	24 25	1 A	v L	G A	L	F. Z	Y K	ET	T S	E G	A G	P. G	K G	V R	F S	C	E	V F	Y L	K E	E D	R F	7 7	F Y	G X	K N	K Q	Y T	V 	W T	F D
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	L C	i 1	G Y	W R	Y A	A M	Z O Z	N S	W S	FS	K F	T	Y G	D V	P S	S G	! H	N V	C V	Ŧ	V D	E	E S	M G	S	E	A M	V A	E W	G F
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	H L	1	TE	TQ	E	l Q	V G	M G	L S	N Y	P K	A K	N	T G	R Y	S Y	- D	s s	7	M K	T D	S	Q L	E	F W	v s	E	K T	LD	T K
SEO. SEO.	ID. ID.	NO. NO.	24 25	K W	R	L	K G	R S	H P	P P	E	E	T	G	G	F	Q	E	A T	PF	L	A F	Y	D S	A Q	I K	W	A F	L I	A S	L V	A S	٧ ٢
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	N L	K S	T	S	G	G	G V	G L	R A	s v	G V	C	R L	L S	E	ロン	F	N Y	Υ N	N S	N H	0 V	F	i Y	T	Δ (I	2.0	s	Y	R P
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	A N	M L	N N	S N	s L	S T	F	E	G G	v C	s s	G L	H A	V L	V A	F	D V	A F	S P	G	S G	R	M D	A G	W Y	T H	L	G	ER	Q S
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	L Q	Q F	G p	G F	s v	Y	K Q	K A	1 R	G L	Y W	Y L	D L	S G	T L	K G	D F	D S	L	S G	W Y	S G	K S	T M	D F	K T	W K	1	G W	G W
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	s V	P H	P T	A V	D F	Q T	ı K	L K	V E	! E	K K	T K	F	R W	F	L K	s T	Q	K E	L P	F W	I K	S L	V Y	S A	V T	L V	S G	S L	L L
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	G V	1 G	V M	L	A V	V L	V T	C L	L A	s	F W	0 7	! !	Y V	Z 0	S P	H	V H	F. F.	Y	1	Q E	N T	SF	Ω Α	P K	N E	L	N P	N K
SEQ. SEQ.	ID. ID.	NO. NO.	24 25	L E	T	A	V D	G V	c s	s	L	A P	L Q	A	A E	V H	F C	p S	L S	G K	L K	D M	G N	Y T	н W	l L	G G	R I	SF	Q Y	FG	P Y	F K
SEQ. SEQ.	ID. ID.	NO. NO.	. 24 . 25	V G	C	QL	A L	R	L	W G	L	L	G	L	G Y	F	S T	L K	G S	Y V	G S	S	M E	F K	T 1	K N	i D	W H	W R	V A	H	T G	V M
SEQ. SEQ.	. ID. . ID.	. NO.	. 24 . 25	F	T I	K Y	K	E	E	K V	K	E	W	' R I	K T	T A	L P	E	P	W	K	L	Y S	A S	T Q	v Q	G	L A	L	V F	G A	M F	D A
SEQ SEQ	. ID.	. NO . NO	. 24 . 25	۷ \$	L	T A	Ł	A	l F	w s	s S	1 Y	\ '	D T	P	L V	H V	R	T	1 Y	EI P	T	F	A	K	E	E	P	K	E	D E	I W	D Q
													F	_	IG	7.	7	b															

SUBSTITUTE SHEET (RULE 26)

SEQ. ID. NO. 25

SEQ. ID. NO. 25

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 SEQ. ID. NO. 24
 V S I L P Q L E H C S S K K M N T W L G I F Y G Y K G L L L SEQ. ID. NO. 25
 S E T Q D T M K T G S S T N N N N E E E K S R L L E K E N R E

 SEQ. ID. NO. 24
 L L G I F L A Y E T K S V S T E K I N D H R A V G M A I Y N SEQ. ID. NO. 25
 L E K I I A E K E E R V S E L R H Q L Q S R Q Q L R S R R H

 SEQ. ID. NO. 24
 V A V L C L I T A P V T M I L S S Q Q D A A F A F A S L A I SEQ. ID. NO. 25
 P P T P P D P S G G L P R G P S E P P D R L S C D G S R V H

 SEQ. ID. NO. 24
 V F S S Y I T L V V L F V P K M R R L I T R G E W Q S E T Q SEQ. ID. NO. 25
 D T M K T G S S T N N N E E E K S R L L E K E N R E L E K I SEQ. ID. NO. 25

SEQ. ID. NO. 24 I A E K E E R V S E L R H Q L Q S R Q Q L R S R R H P P T P

SEQ. ID. NO. 24 PDPSGGLPRGPSEPPDRLSCDGSRVHLLYK

FIG. 7c.

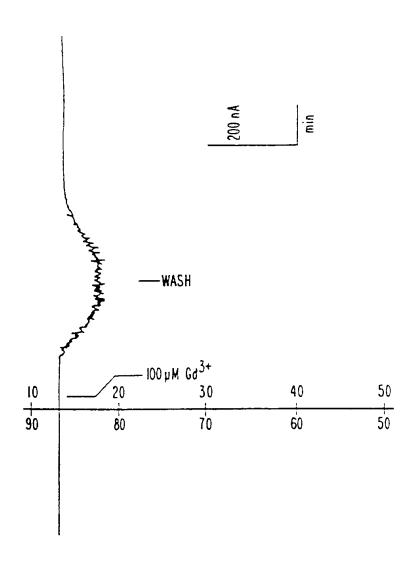


FIG. 8.

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ClustalW Formatted Alignments

SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	A T G G G A T C G C A T G G C A T T T T	A G G G A A A G C G A T C A G T G C T T G C G C T C C C G G A T A G C T G C T G C T G G T G C T T G C G C T C C T G G
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	C A C T G C T G C T T C C T C T T G G C	T T G T T T C T T C C T C T T G C T G T G G G G T G C T G T A C T C A C C T G G C A C A C G C T G T G G G G T G C T G T
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	G	T T C T A C T G G A T C C T C C C A G C C A A G A A G G T G G A C C A G C G A C C A G C G A C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A G C C A A G C C A A G C C A A G C C A A G C C A A G C C A A G C C A A G C C A A G C C A A G C C A A G C C A A G C C C A A G C C A A G C C A A G C C C A G C C C A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A G C C C A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A G C C C A A A C C C C
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	C T G A C C C T G G G C C C A A A A G A	A A A G A A C T C A C A G C C A G G G A G A C T T G G T G C A G G G G A C A T T A T C C A G G G G A G A C T T G G T G C
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	T G G G T G G G C T T T G G G G G G C T	C C A T T C C A T A C G G G T G T T C C A G T G C A C C A C T T T C C T A T T C A T T T G T T C C C A G T G C A C C A
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	G	A T T A T T T T G G G G G G T C C A G C A G A G A G G A C T G T G C A A G A G A C T G T C A A G A T C A G C A G C A G A G A C T G T
SEQ. ID. NO. 34 SEQ. ID. NO. 30	G	T C C A C G C A A A G G G A G A T G A G C A C C G T G G C A G G C C G G A G T C T G T G G A T G A G C A C C G T G G C A
SEQ. ID. NO. 34 SEQ. ID. NO. 30	T C C A G C G C C T A A T G T A T C A G	G C C T T G T G G G G A G C T G G G A G C T T T T G G G G A G C T T T T G G G A G C T T T T C C G T G G G G A G C C A T G C T T T T T C C G A G G C C A T G C T T T T T C C G A G G C C A T G C T T T T T C C G A G G C C A T G C T T T T T C C G A G G C C A T G C T T T T T C C G T G C T T T T T C C G A G G C C A T G C T T T T T C C G A G G C C A T G C T T T T T C C G A G G C C A T G C T T T T T C C G A G G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G A G C C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T C C G T G C A T G C T T T T T T C C G T C A T G C T T T T T T C C G T T T T T C C G T C A T G C T T T T T T C C G T C A T G C T T T T T T C C G T C A T G C T T T T T T C C G T C A T G C T T T T T T C C G T T T T T T C C G T T T T

SEQ SEQ SEQ	. ID . ID . ID	. NO . NO . NO	. 34 . 30 . 26	G	G	C	A T A	CCC	G T	G G	G G	A G A	C C	C C	G G	C C	A C A	T A T	C G C	A G A	A C A	C	A C	G T G	T G T	A G	A T A	
SEQ. SEQ. SEQ.	. ID. . ID.	. N O.	. 34 . 30	T	C T	G T	C G	A C	C C	C A	T T	G A	C G	T A	G G	C G	C A	T G	G A	G T	C A	G A	T A	G C	C A	G	T C C C	C A
SEQ. SEQ. SEQ.	ID. ID.	NO.	34 30	T G	G C	G C	G C	T A	G G	C	A C	C C	A T	C T	A C	T T	C T	C C	T C	C C	G A	A A	C	A T	G T	T G	C T A T	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	C G	T C	C T	C G	A G	A G	G A	G T	A A	C C	A A	C G	A G	C A	A T	T A	G T	C T	G T	C G	T A	G C	G A	T A C A	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	A G	G C	G A	C A	A C	C A	T C	G C	G G	A T	C T	T T	T C	T T	G A	T A	G G	C G	G C	T C	G T	C T	G C C	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C A	A A	C G	T C	C	A A	G C	C C	C C	G T	T G	G A	G G	T T	G T	C T	T T	G G	A T	T T	G G	G C	C T	C T C T	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	A A	C A	G A	C C	C A	A A	C A	A A	T T	C T	T G	G A	C T	C T	C C	C T	G T	A T	C G	G A	G A	C C	T C	C T	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T G	A A	T T	G G	C A	G G	A T	C T	C C	C T	A G	T C	G A	G A	T C	G T	A G	T C	G T	C C	T A	C G	C A	C G	A C

FIG. 9b.

SEQ. SEQ. SEQ.	. ID. . ID.	NO.	34 30	C A	T C	G A	C T	C T	A C	T C	C C	A T	C	T T	G A	G C	T G	G A	T T	T T	A G	T C	T T	G G	G T	C G	A G G	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	T G	T G	C G	C A	T G	A C	C A	A A	G C	T T	G G	A G	T C	G T	C	C A	T G	C G	C	A G	T T	C C	C T	T A C A	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	G A	T C	G G	G G	C C	C A	A G	A T	C G	C G	T C	C A	T A	T A	G T	A C	G T	G G	C C	T T	A G	T G	T G	G T G T	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	A T	G C	A T	T T	C C	C T	C A	A C	C A	A T	G T	A C	T C	T C	A C	G A	C G	T G	A T	C C	G A	C	C T	A T T	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T T	A G	C	C	A T	G C	T C	G T	C C	C C	A A	A G	G C	C A	T G	G A	A C	G T	T C	G C	A T	C C	A A	G A G A	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T A	C A	C C	C A	G A	C G	T A	A A	T T	G C	A A	C A	T T	A T	C C	T A	T A	T G	G T	C C	C T	C T	G T	A C C C	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	A C	G C	T G	G A	C A	C C	T C	C A	C T	T C	G C	A C	C C	T A	T A	C T	T G	T A	C T	C G	A A	A G	C	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C	A C	A A	G G	G G	C C	C C	A A	T C	G T	G G	C C	T C	G A	A T	G	A G	T C	T A	C G	T A	C C	C A	G T	C C

FIG. 9c.

SEQ. SEQ. SEQ.	. ID. . ID.	NO NO	. <i>34</i> . <i>30</i>	T A	T T	C	T G	T A	C G	A T	A A	C T	T	G T	G C	A C	C G	C	T T	A G	T G	G A	T A	G C	T T	C G	C C C	G
SEQ. SEQ. SEQ.	ID. ID.	NO.	. <i>34</i> . <i>30</i>	C T	T G	G G	T G	G C	G A	C	G A	T A	C T	T T	G G	A C	G A	G G	G C	C T	G G	A A	C T	T G	A A	T C	T G G G	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	34 30	C	G T	A A	G T	A G	C G	A G	G C	G G	C G	A C	T C	T G	G G	A G	G G	G A	C T	C T	T G	T A	T G	G A	G A A	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	T T	A C	G C	A G	G A	G G	C A	T G	C G	G A	Τ Λ	G G	C C	C T	C	G A	C G	A G	A A	C A	A A	T G	0000	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	G A	T T	G A	T T	G C	G T	C G	C C	A A	C T	C	T G	C A	G C	G T	A T	G C	A A	A G	A T	G G	T A	G A	A G C G	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C	C A	G T	T C	G T	C C	C C	A C	T A	G G	A T	G A	C C	C T	G C	C T	G G	C A	G T	G G	C A	C G	T G	C T A T	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	G	A A	G G	G A	G T	T C	G C	T A	G G	G C	T A	G T	C G	G T	A G	G G	C T	C A	C G	T A	G G	C G	T T	G G	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C A T A	G T	A C	A A	G A	C A	C A	C T	A T	G C	T	G A	C C	C G	C	G C	C	G A	T A	G A	G G	C T	T	G A	T T

FIG. 9d.

SEQ SEQ SEQ	. ID. . ID.	NO.	. 34 . 30	C	C G	T T	G G	T G	T T	C T	A T	C T	C C	C T	G C	T	T A	C G	T T	G G	A G	G C	G C	A C	T A	G G	C A C	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	34 30	C C	G T	G T	G G	A A	G G	C C	T C	G C	C	T T	T C	G A	C T	T C	G A	C A	C G	A G	G A	C G	C A	A T	T G T G	C G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G T	C C	C C	T G	C G	A C	A G	T C	G A	C A	C T	A A	G T	C C	T A	T C	C G	A G	C G	C C	T A	G A	G G	G G A G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	G C	G T	C G	C G	A C	G T	T G	G G	A C	T C	G A	G G	T C	T G	G A	G G	G G	G C	G C	G T	C G	C G	C G	G T C T	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G A	A G	G C	A T	G C	T C	G T	T C	G C	G C	T T	G G	G A	C T	A C	G G	G C	C C	A A	G T	T G	G C	A C	A G T G	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G A	G G	G T	C A	T C	G T	C T	T C	G C	A A	G C	G G	G T	T G	G G	C T	T T	A G	T G	C C	A G	C G	C C	C A A	T C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C C	G A	A T	G T	C G	T G	G A	G T	C T	C C	T G	C C	C T	T C	A T	C G	C A	C A	C G	A G	T C	C T	A G	G	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	G C	A A	C G	T A	T T	T C	G C	C C	C A	T G	C G	C C	T T	A T	C	T C	T G	C G	C G	A A	G A	A T	G T	C C	C C

FIG. 9e.

SEQ SEQ SEQ	. ID. . ID.	. NO . NO	. 34 . 30	T T	G	G A	A A	C G	C A	C A	T G	T G	G	G C	A C	A A	C T	A C	A C	C	A A	G G	C G	C A	G A	G G	A T	A C A
SEQ. SEQ. SEQ.	. ID. . ID.	NO.	34 30	C T	C	G C T C	C C	T C	G A	G C	T A	T A	C T	C	G G	T T	G T	A T	A T	T G	T C	C C	T A	G A	G G	G	A A	G G
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	A T	C G T G	A T	G G	G G	T G	T A	C A	C G	G A	C A	T A	G C	C A	A T	G T	C T	T A	T A	C C	C T	G G	G C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	A A	G C	G C C C	G T	A C	G C	A A	C A	T G	G A	C A	G G	C G	A T	G G	C C	C A	C A	A A	C A	T G	C G	T A	C C	T C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	C T	G G T G	G A	G C	C C	T T	G G	T T	G G	C G	C A	C C	T A	T C	T C	G T	A T	G T	C C	A T	G	G	A G	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T G	C G	A C T C	A C	A A	G C	A G	T A	C A	A G	T A	G A	T A	T G	T T	G G	T G	G C	G G	T A	C C	A A	A G	T G	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	A T	G A	T G	G C	T A	A A	C C	G A	C G	C C	A T	T C	G G	G A	C C	C A	C G	A C	T C	G T	C T	G C	C C	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C A	C	A C	C	A C	A T	C C	A T	T G	G T	C A	A C	C A	C G	G G	T G	G G	C A	C T	C G	T A	C G	T A	G A	C C

FIG. 9f.

SEQ SEQ SEQ	. ID. . ID.	NO.	. 34 . 30	C A	C T	C	A A	A G	C	A A	C G	C T	A G	C T	C	C G	G A	G G	C A	T C	C	T C	G C	T T	G T	A A	T C C C	G A
SEQ. SEQ. SEQ.	. ID. . ID.	NO. NO.	34 30	C T	G A	A G	T A	G T	C T	G A	G C	C A	C C	A G	G C	T A	T T	A T	A T	C A	G C	G G	G G	C A	G T	C A	C C T C	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	C C	C T	T A	C C	T A	A A	C T	A G	A T	G G	G T	A A	C C	T T	T T	T A	G G	T C	G A	C G	T T	C C	A T	C A A	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G T	T C	C C	A A	A T	G T	T G	T C	T C	G C	A A	T C	G G	C C	C C	C T	C T	C G	T C	T A	T A	C G	G A	A C T C	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	A A	G T	C A	T T	G A	A C	C C	A T	C G	C C	C T	A T	C A	A C	A C	T T	G G	A G	G G	G A	T G	C A	C C G C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C G	T C	T T	T C	G T	A T	C C	C A	C	C C	T A	T A	T T	G G	G G	T C	G T	A C	T C	G T	G G	T T	A G	A T C T	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G G	G A	C	C A	G T	C	T A	A A	C G	A A	A A	C A	A G	T T	C T	T G	T A	C G	A G	C	C G	T T	A G	G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T A	G G	C G	G T	T C	G C	C T	A G	G A	G A	C G	A C	G A	T C	G C	G T	G A	C C	G G	C G	T C	A A	T T	G C C	G T

FIG. 9g.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	. 34 . 30	C A	T A	A A	C C	C T	A T	G T	A A	A C	G A	G A	T A	G C	G A	G A	C T	T A	A T	C G	G	G	G C	G G	Α	Α.
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G C	A A	A G	G G	G T	C G	T A	T C	G C	A T	C T	T T	C G	T A	G T	G G	A A	C G	A T	C G	C T	A G	G G	C T	
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T A	C C	A C	T T	C G	C G	C T	A G	T G	G G	G G	G A	C A	C	T T	C A	A T	C T	C C	C C	T .÷.	C T	Λ C	A G A G	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C C	G A	G A	C C	C T	C G	C G	C C	T A	G C	C C	C T	C C	G T	C C	C C	T C	C C	T A	C G	G A	C G	T G	A G A G	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	A G	G G	T C	G T	A C	G C	C A	C T	C C	T G	G T	C G	C T	T T	C T	C A	A A	G G	A G	A A	T A	G G	A T	A G C G	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T G	G G	A T	A A	G T	A T	G A	T C	G A	T A	G C	C G	A T	G C	C T	C A	G T	G G	G C	C C	G A	A A	A G	A G A G	T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C G	T G	G	C A	T G	G A	C A	T A	G G	G A	C C	T T	С	T T	G T	C C	A A	T T	T C	C A	C A	G C	T G	G A	C G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C G	A A	G	C A	C A	C A	T A	A T	T C	G C	A T	G G	T T	A G	C G	C A	G	A T	T G	T G	G G	G T	A T	C C	G T

FIG. 9h.

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SEC SEC SEC). ID). ID). NC). 34). 30	A C	. A	T A T	G	C G	A G	C A	T	G	G T	C	G	C	T	G T	A T	T	T	G	T	G A	G A	C	C T	T G
SEQ SEQ SEQ	. ID . ID	. NC . NC	. 34 . 30	G C	G A	000	C	T C	A G	C A	T	G A	G C	C T	C G	C	A C	A T	T G	G G	C	C A	A G	G	C G	C A	T C	G C
SEQ SEQ SEQ SEQ	. ID. . ID.	. NO . NO	. 34 . 30	A A	C G	G T G T	G A	G A	C A	T G	G G	C G	T A	T T	C C	G A	A T	A T	C G	T A	G G	C G	C G	C G	C G	A A	G G	G C
SEQ. SEQ. SEQ. SEQ.	. ID. . ID.	. NO.	34	A C	G C	C T A T	A C	C C	A T	T G	C C	C T	G G	C C	T T	G T	G T	G G	G A	C G	G T	A G	T T	G G	C T	C G	T G	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	34 30	G G	G T	G C G C	T T	G C	T C	G T	G G	G A	A T	C G	C G	T G	G G	T A	C G	A T	C A	C T	A A	T G	C T	G G	C A	.L C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T G	G A	A C G C	C A	T C	C A	G G	G A	T T	G G	C C	C C	C A	T G	G T	G G	C C	C C	A T	C G	C T	C A	T A	C C	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	T A	T	G T	T G	G C	C C	T C	G A	G G	G A	T T	G G	T A	C C	T T	T T	T C	G T	T G	G G	C T	G C	G C	C A	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	A G	T A A	T G	G A	C A	C C	A C	C A	A C	C A	C C	A C	G T	T C	G C	G T	T G	C C	A T	A T	G C	G G	C A	C A

FIG. 91.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	T C	C T	A G	G C	G C	T C	C C	G A	0000	G G	A A	G	C T	T A	C	T A	G T	C C	T C	A G	C	A T	T G	C G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T G	G C	C G	T A	G T	G G	G C	T C	C G T G	G G	T G	G G	T C	C T	T G	T T	C G	C G	T G	C A	T C	G C	C T	T G	A T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	C C	T A	G C	C C	A A	T T	G C	A G	A C C C	C	T T	T G	C	A C	T T	C	T G	T G	C T	A G	T C	T C	G C	C T	C G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	A G	A C	G C	C A	C C	A C	T C	C T	0 0 0	A T	C T	G T	G G	C T	A G	G C	T T	G G	T G	G G	T T	A G	C T	C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	T T	A T	C G	G T	G G	C C	G G	T G	T C C C	T A	T C	G A	G A	T T	T G	T C	G C	G A	G C	C A	A C	C C	T A	G G	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C G	T G	T T	C C	T A	C A	G	G G	C T C T	C C	T T	C C	C A	T G	A G	C T	T C	C G	A G	G G	C A	C G	C C	T T	C C
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	C T	T G	C C	A T	C A	C C	A A	A T	С	A C	C T	C G	A C	A T	C G	C G	G G	C T	A G	T G	T T	G	C T	A C	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>	G T	C	A C	T T	C C	T T	T G	C	G T	G A	T C	G T	G G	G C	G A	C T	C G	C A	G C	G C	G T	A T	G C	G A	G T

FIG. 9j.

SEQ SEQ SEQ	. ID . ID:	. NO . NO	. 34 . 30	T C	G T	C	C	C A	A T	G T	C	Ģ C	C	C A	C	A G	. C	G C	C A	T T	C	C	A A	T	C	A G	C	A T A T
SEQ. SEQ. SEQ.	. ID. . ID.	NO.	34 30	C G	C T	T G	G T	C G	C T	T A	C	A C	C T	A T	G A	G C	T G	G	G C	C G	C T	A C	T T	C T	T G	G	C T	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T T	G G	G G	C G	A C	C A	T C	T T	A G	Т С	C C	T T	C T	G C	G T	G C	C T	C	A T	G C	C T	T	G C	C C T C	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C C	A T	T C	C A	G G	T C	G C	G C	T T	C G	G C	C T	C C	T A	G C	G C	C A	T A	G G	G A	T C	G C	G A	T T A T	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G C	A G	G C	G A	C T	A T	C G	C C	G A	G C	G G	C C	A A	C T	A C	G T	G T	C C	A G	A G	G T	G G	A G	T G G	A G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C C	A C	G C	C G	C G	C G	C A	C G	G G	A G	A T	C G	G C	G C	C C	G A	G G	G C	A G	G G	G C	T C	G A	G G C G	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G C	A T	C T	A C	C A	T T	G C	C A	G G	C T	T C	G C	C T	A G	A C	C	C T	A C	C A	C	G A	C G	G	A T	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G G	C C	A C	A A	G T	T C	A T	T G	G C	T C	T T	G G	G G	C	C A	T C	C T	G T	C A	T T	G C	G T	C C	С	T G

FIG. 9k.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	A G	C C	A A C A	A A	T G	C C	T T	G G	C	T T	C	C A	T T	C	A G	T T	C	G G	C T	C	C G	T C	C	T T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C G	A C	A C T C	G G	C G	T T	T G	T G	A T	T G	G G	C A	C G	T G	T C	C A	A C	A C	G G	A G	C G	T C	C A	G C	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	A G	A G	T G C G	T A	G A	C G	C G	C A	C G	G A	A C	A A	A G	A C	C C	T C	T C	C	A G	A A	C A	G C	A G	G G	G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C	C G	A A G A	A A	G G	T G	T T	C	A G	T T	T G	G A	G C	C A	T C	T T	C G	A C	C G	C C	A T	T G	G C	T A	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C C	A C	T C A C	C C	A C	C G	C C	T G	G A	C T	A G	T C	C A	A A	T G	C T	T A	G T	G	C T	T T	G G	G G	C G	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	T T	T C	T C G	C C	T T	G G	C G	C C	C C	A T	T A	C C	T A	T A	C T	T G	A T	T G	G C	T T	C C	A C	C T	C C	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C T	C C		G C	T G	G C	A T	C C	T T	A G	C	C A	G C	G G	G C	T T	A T	C T	A A	G T	A G	C C	C C	A T	C T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C C	A A	C A	C T	A A	T C	G T	T C	G	C C	G A	T A	G G	T T	C	A C	G C	T C	C C	A G	G A	C A	C A	T A	C C

FIG. 9L.

SUBSTITUTE SHEET (RULE 26)

SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	A G C G	GCTC	C G T G G G G C C A	T G C T T G A G T T C A	C A G C A G G C T G C C T T G G C T G C T G C C
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	TCTT	T G C G C A T G	C C C A A T A C A C	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	A A C A A C C A T C A T C A T C A T C A T C A T
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	CCTC	T T C C C T G G	A G C C G C A T T G	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	T T A A G T A A C A C C A T C T T C A A C G T G
SEO. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	A T C G T A T G	A G G A T C A C	G G T G C C T C C A	G T T G C A G T G A C T	T G C T C T G C A C C G G C C A G C C
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	C A G C T A C A	T C A C G A C C	G C T T T A C C A C	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	T A T A A T G G C T G C C A G G G C C
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	C C G G A G T C	G C C A A G C C	C G C T G T C A G C	C G C C G C G C G C T C C	A C C A T C A G C A A C G T G G T G G G G T C T
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	G T C T C T T G	C C C G G C T G	C A A G C C C T C T	G G T C C A T T G C G C	CCAAGC
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	T T G G T G C A	A G G C C A T C	T C C A C A T C C T	G G G A T C C T T C C A	C A C C C C A

FIG. 9m.

SEQ SEQ SEQ SEQ	. ID.	. NO . NO	. 34 . 30	C G	T A	C C A C	C G	T A	C A	C	T G	C T	C	A G	T T	C T	A A	G	C	A C	G	C	А	Α	G	Α	G	C
SEQ. SEQ. SEQ.	. ID. . ID.	NO NO	. 34 . 30	Α	Α	c c c	Α	G	C	G	Α	Α	G	Α	C	С	С	Α	T	T	С	С	C	Α	С	Α	G	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	. 34 . 30	С	С	G G G	Α	G	Α	G	G	С	Α	G	Α	Α	G	С	Α	G	С	Α	G	C	Α	G	C	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G	С	C T A	G	G	С	C	С	T	Α	Α	C	C	C	Α	G	С	Α	Α	G	Α	G	С	Α	G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30		Α	C G C	C	Α	G	С	Α	G	С	С	С	С	T	G	Α	С	С	С	T	C	С	C	Α	С
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	Α	G	A C G	Α	G	С	Α	Α	С	G	Α	T	С	T	С	Α	G	С	Α	G	C	Α	G	С	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	G C T	G A	C G	C A	C T	T G	A C	A A	C A	C G	C C	A A	G G	C A	A A	A G	G G	A T	G C	C A	A T	G C	C T	A T	G T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30	C		G C																						

FIG. 9n.

SEQ. SEQ. SEQ.	. ID. . ID.	NO.	34 30													G A
SEQ. SEQ. SEQ. SEQ.	. ID. . ID.	NO. NO.	34 30													
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30													A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>													C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30												A G	
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30												A C	
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	34 30													
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	<i>34</i> <i>30</i>													

FIG. 90.

SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	G A A G T G	A C G G A G	G A C	TTAC	G A T C G A A G	T G A C A G T T	C G T C C G T C C C
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	A G G C A G	A A A C A C	C A G T T G	G T C T	T G C A T G T C	AGGA CAGT	C C T G T T C A C A
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	G G G G A G	T G G C T T	A G A T G T	C C A C	G C G G	C C A G G G T G	A G G T G G A G G C
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26							
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26			TAG	тстс	CCAG	ТТСА	CAGAG
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	СТТ	TGT	CAT	CAGT	rgg T (G G A G	GCAGC
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	АСТ	G T T	АСА	GAAA	A A C G	TAGT	GAATT
SEQ. ID. NO. 38 SEQ. ID. NO. 34 SEQ. ID. NO. 30 SEQ. ID. NO. 26	СА						

FIG. 9p.

ClustalW Formatted Alignments

SEQ. II SEQ. II SEQ. II	D. NO. D. NO.	. 35 . 31	M M	G A	S F	L Y	L S	A C	L C	P W	A V	L L	L L	L A	L L	W T	G W	A H	V T	A S	E A	G Y	P G	A P	K D	I K Q K	V R
SEQ. IL SEQ. IL SEQ. IL	D. NO. D. NO.	35 31	L A	T Q	L K	E K	G G	D D	L I	V I	L L	G G	G G	L L	F F	P P	٧ I	H H	Q F	K G	G V	G A	P A	A K	E D	GPQP	C D
SEQ. IL SEQ. IL SEQ. IL	D. NO. D. NO.	35 31	L	P K	ν \$	N R	E P	H E	R S	G V	I E	Q C	R I	L R	E Y	A N	M F	L R	F G	A F	L R	D W	R L	I Q	N A	H R M R	D I
SEQ. IL SEQ. IL SEQ. IL	D. NO. D. NO.	35 31	P F	H A	L I	L E	P E	G I	V N	R S	L S	G P	A A	H L	I L	L P	D N	S L	C T	S L	K G	D Y	T R	H I	A F	G L D L	E T
SEQ. II. SEQ. II. SEQ. II.	D. NO. D. NO.	35 31	Q C	A N	L T	D V	F S	v K	R A	A L	S E	L A	S T	R L	G S	A F	D V	G A	S Q	R N	H K	I I	C D	P S	D L	I G N G	S L
SEQ. II SEQ. II SEQ. II	D. NO. D. NO.	35 31	Y D	A E	T F	H C	G N	D C	A S	P E	T H	A I	I P	T S	G T	v 1	I A	G V	G V	S G	Y A	S T	D G	V S	S G		Q S
SEQ. II SEQ. II SEQ. II	D. NO. D. NO.	35 31	V T	A A	N V	L A	L N	R L	L L	F G	Q L	I F	P Y	Q I	I P	S Q	Y V	A S	S Y	T A	s s	A S	K S	L R	S L	D L	K S
SEQ. IL SEQ. IL SEQ. IL SEQ. IL	D. NO. D. NO.	35 31	S N	R K	Y N	D Q	Y F	F K	A S	R F	T L	V R	P T	P I	D P	F N	F D	Q E	A H	K Q	A A	M T	A A	E M	I A	L D	R I

FIG. 10a.

SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	FFNWT	Y V S T W N W V	V A S E G D Y G T I A A D D	Y V S T L A S E G G E T G I E A F E D Y G R P G I E K G E T G I E A F E
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	L E A R A F R E E A	R N I C E E R D	V A T S E K V I C I D F S E	I G G V C I A Q S G R A M S R A A F L I S Q Y S D E E G R A M S R A A F
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	EGVVR	A L L Q V E V I	K P S A R V A Q N S T A K V	K R L L E T P N A V L F T R S E D A I V V F S S G P D V L F T R S E D A
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	R E L L A L E P L I	ASQR KEIV	LNASFTWRRNITGK	E A A K K L N Q S V A S D G W G A L I W L A S E A W A V A S D G W G A L
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	ESVVA SSSLI	G S E G A M P Q	A A E G A I T Y F H V V G G	P V Y Q Q E E I A I E L A S Y P I S T I G F A L K A G I E L A S Y P I S
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	D F A S Y O 1 P G F	FQSL REFL	D P W N N S R K K V H P R K	D R Y F R S R T L N P W F R E F W E S V H N G F A K E N P W F R E F W E
SEQ. ID. NO. 35 SEQ. ID. NO. 31	Q R F R C F W E E T	S F R Q F N C H	R D C A A H S L Q E G A K G	F G C K L G S H G L R A V P F E Q E F L P V D T F L R L R A V P F E Q E
SEQ. ID. NO. 35 SEQ. ID. NO. 31	S K I M F G H E E S	V V N A G D R F	V Y A M A H A S N S S T A F	R D S S Y E Q E G L H N M H R A L C R P L C T G D E N L H N M H R A L C

FIG. 10b.

SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	PNTTF	LCDA	AMRPVNG IDYTHLR	L H N M H K I R R L Y K D F I S Y N V Y I R R L Y K D F	F V L N L A V Y
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	V K F D A	PFRF	P A D T H N E I Y T C L P G	ELLGYIF VRFDRFC RGLFTNC VRFDRFC	D G I S S C A
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	GRYN I	FTYL EAWQ	L R A G S G R Q V L K H L R	DAPGRYE YRYQKVO HLNFTNN YRYQKVO	Y W A I M G E
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	EGLTI	D T S L	LIPWASP	W T N Q L H L S A G P L P A I I N W H L S S A G P L A A	SPED
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	SEPCI	QNEV	V K S V Q P G G Y Y N V Y A	C S L P C K F E V C C W L C K K G E R L F E V C C W L C	I P C F I N E
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	Q P Y E Y E K I L V	R L D E V S G F S	EFTCADC SREVPFS	C E G Y N Y C G L G Y W P N N C S R D C I G L G Y W P N	ASL AGT
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	TGCFI	ELPQI IEGEI	E Y I R W G D P T C C F E C	AWAVGPY VECPDGI	V T I A E Y S D
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	C L G A E T D A	L A T L I S A C N I	F	RHNATP' SNENHT	V V K A S C F E

FIG. 10c.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	35 31	S	G	R	E	L	C !	Y R	l W	L G	D L D L	G A	G W	V A	F V	L G	C P	Y V	C	M I	T A	F C	i L	F G	I A	A L
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	35 31	K A	P T	S L	T F	A V	V L	C G	T V	L F	I R V R	R R	L H	G N	L A	G T	T P	A V	F V	S K	V A	C S	Y G	S R	A E	L L
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	35 31	L C	T Y	K I	T L	N L	R G	I G	A V	R F	C I L I	F C	G Y	G C	A M	R T	E F	G I	A F	Q I	R A	P K	R P	F S	I T	S A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	35 31	P V	A C	S T	Q L	V R	A R	I L	C	L L	T A G A	L T	I A	S F	G S	Q V	L	L Y	S	V A	V L	A L	W T	L K	V T	V N
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	35 31	E R	A I	P A	G R	T I	G F	K G	E G	T A	V A R A	P E	E G	R A	R Q	E R	V P	V R	T F	L I	R S	C P	N A	H S	R Q	D V
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	35 31	A A	2 1	M C	L L	G A	S L	L I	A S	Y G	K N Q N	V L	L L	L I	I V	A V	L A	C W	T L	L V	Y V	A E	F A	K P	T G	R T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	35 31	K G	C K	P E	E T	N A	F P	N E	E R	A R	K E	F V	I V	G T	F L	T R	M C	Y N	T H	T R	C D	I A	I S	W M	L L	A G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	35 31	F S	L L	P A	I Y	F N	Y V	V L V	T L T	2 1 2	S A	D L D	Y C Y	R T R	V L V	Q Y	T A	T F	T N	M T	C R	V K	S C	V P	S E	L N

FIG. 10d.

SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	SGS	V V L A K F	G C L I G F	F A P T M Y	I Q T T K L H I T T C I K L H I	I L F I W L	Q P Q I A L L I	K N T P I F
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	I E E	VRC	STA	ана	V Y I I F K V A T M C V	ARA	TLRI	RSN
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	VSR	KRS	S S L	GGS	K V A A T G S T L F Q P	P S S	SISS	SKS
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	NSE	DPF	P Q P	ERQ	GSTP KQQQ SSSL	PLA	LTQ	QEQ
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	QQQ	PLT	L P Q	QQR	Q Q Q P S Q Q Q T T S S	PRC	T Q Q I K Q K '	E Q Q V I F
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27	Q Q P G S G	L T L T V T	P Q Q F S L	Q R S S F D	QQQP EPQK	R C K N A M	QKV AHG1	I F G N S T
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27								
SEQ. ID. NO. 39 SEQ. ID. NO. 35 SEQ. ID. NO. 31 SEQ. ID. NO. 27								

FIG. 10e.

```
SEQ. ID. NO. 39 ETDLDLTVQETGLQGPVGGDQRPEV
SEQ. ID. NO. 35 VEDPEELSPALVVSSSQSFVISGGG
SEQ. ID. NO. 31
SEQ. ID. NO. 39 EDPEELSPALVVSSSSQSFVISGGG
SEQ. ID. NO. 35 STVTENVVNS
SEQ. ID. NO. 31
SEQ. ID. NO. 37
SEQ. ID. NO. 37
SEQ. ID. NO. 35
SEQ. ID. NO. 31
```

FIG. 10f.

ClustalW Formatted Alignments

SEQ. IC SEQ. IC SEQ. IC SEQ. IC	D. NO. D. NO.	46 36	A A	T T	G G	G G	G G	A A	T T	C C	G G	C C	T T	G G	C	T T	T T	G G	С	G	C	T T	C C	C C	C	A G G G	G G
SEQ. IC SEQ. IC SEQ. IC SEQ. IC), NO.), NO.	46 36	C C	A A	C C	T	G G	C C	T T	G G	C C	T T	G G	C C	T T	G G	T T	G G	G G	G G	G G	T T	G G	C C	T T	T G G A	T T
SEQ. IE SEQ. IE SEQ. IE SEQ. IE), NO.), NO.	46 36	G G	G G	C C	T T	G G	A A	G G	G G	G G	C C	C C	C C	A A	G G	C C	C C	A A	A A	G G	A A	A A	G G	G G	T T T G	G
SEQ. IE SEQ. IE SEQ. IE SEQ. IE). NO.). NO.	46 36	C C	T T	G G	A A	C C	C C	C C	T T	G G	G G	A A	G G	G	G G	A A	G G	A A	C C	T T	T T	G G	G G	T T	C G G C	C C
SEQ. IC SEQ. IC SEQ. IC SEQ. IC). NO.). NO.	46 36	T T	G G	G G	G G	T T	G G	G G	G G	C C	T T	G G	T T	T T	C C	C C	C C	A A	G G	T T	G	C	A A	C C	G C C T	A A
SEQ. IL SEQ. IL SEQ. IL SEQ. IL	D. NO. D. NO.	46 36	G G	A A	A A	G G	G G	G G	C C	G G	G G	C C	C C	C C	A A	G G	C C	A A	G G	A A	G G	G G	A A	C C	T T	G G G A	T T
SEQ. IE SEQ. IE SEQ. IE SEQ. IE	D. NO. D. NO.	46 36	G G	G G	T T	C C	C C	T T	G G	T T	C C	A A	A A	T T	G G	A A	G G	C C	A A	C C	C C	G G	T T	G G	G G	C C	A A
SEQ. IL SEQ. IL SEQ. IL SEQ. IL	D. NO. D. NO.	46 36	T T	C	C C	A A	G G	C C	G G	C C	C	T T	G G	G G	A A	G G	G G	C C	C	A A	T T	G G	C	T	T T	T T	T T

FIG. 11a.

SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T	G G	A C C T	A A	C C	T T	G G	G	A A	C C	C	G G	C	A A	T T	C	A A	A A	C	C	G G	T T	G G	A A	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	С	C C	G G G T	C C	A A	C C	C C	T T	G G	C C	T T	G G	C C	C C	T T	G G	G G	C C	G G	T T	G G	C	G G	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	G G	CGGC	G G	T T	G G	C C	A A	C C	A A	C C	A A	T T	C C	C C	T T	C C	G G	A A	C C	A A	G G	T T	T T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	T T	T C C T	C C	A A	A A	G G	G G	A A	C C	A A	C C	A A	C C	A A	T T	G G	C C	G G	C	T T	G G	G G	A A	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	C G G C	G G	C C	A A	C C	T T	G G	G G	A A	C C	T T	T T	T T	G G	T T	G G	C C	G G	T T	G G	C C	C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	A C C G	T T	C C	A A	G G	C C	C C	G G	T T	G G	G G	T T	G G	C C	T T	G G	A A	T T	G G	G G	C	T T	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	C C	G G	C C	C C	A A	C C	A A	T T	C C	T T	G G	C C	C C	C C	C C	G G	A A	C C	G G	G G	C C	T T	C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	A A	T T	G G	C	G G	A A	C C	C C	C C	A A	T T	G G	G G	T T	G G	A A	T T	G G	C C	T T	C C	C	C C	A A

FIG. 11b.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C	T	G	C	C	A	T	C	A A	C	T T	G	G G	T T	G	T T	T T	A A	T T	T T	G	G G	C C	A G G G	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	T T	C	C C	T T	A A	C C	A A	G G	T T	G G	A A	T T	G G	T T	C C	T T	C C	C C	A A	T T	C C	C C	T A A C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G	T	G G	G	C	C C	A A	A A	C C	C C	T T	C	T T	T T	G G	A A	G G	G G	C	T T	A A	T T	T	G T T G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	. 46 . 36	A A	G G	A A	T T	C C	C C	C C	A A	C C	A A	G G	A A	T T	T T	A A	G G	C	T T	A A	C C	G G	C C	С	A T T	C C
SEQ. SEQ. SEQ.	. ID. . ID.	NO. NO.	. 46 . 36	T T	A A	C C	C C	A A	G G	T T	G G	C	C C	A A	A A	G G	C C	T T	G G	A A	G G	T T	G G	A A	C C	A A	G A A G	G G
SEQ. SEQ. SEQ.	. ID. . ID.	. NO . NO	. 46 . 36	T T	C	C	C	G G	C C	T T	A A	T T	G G	A A	C	T T	A A	C	T T	T T	T T	G G	C	С	C	G	A C C C	A A
SEQ SEQ SEQ	. ID	. NO	46	C	A A	G	T	G	C	C	T T	C	C	T	G	A A	C	T T	T T	С	T T	T T	C	C	A A	A A	G	C
SEC SEC SEC). ID	NO NO). 46). 36	C	: A	. A	G	G	C		: A	T T	. G	; G	C	T	. G	A A	G	A A	T	T	C	7	. C		G	C

FIG. 11c.

SEQ. ID. N SEQ. ID. N SEQ. ID. N SEQ. ID. N	O. 46 O. 36	T A T T A T	C	T T	T T	C A	\ A \ A	C	T T	G G	G G	A A	C C	C	T T	A A	T T	G	T T	G G	7 7	C	C C	A A
SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO	O. 46 O. 36	T C C T C T	G	T T	G (G (C G	T T	C C	T T	G G	A A	G G	G G	G G	C C	G G	A A	C C	T T	A A	T T	G G	G G
SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO	O. 46 O. 36	T G C G C G C T	A A	G G	A (C A	A G	G G	C C	A A	T T	T T	G G	A A	G G	G G	C	C	T T	T T	T T	G G	A A	G G
SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO	O. 46 O. 36	A A C T C T T T	A A	G G	A (G C	; c	T T	C C	G G	T T	G G	C C	C C	C C	G G	C C	A A	A A	C C	A A	T T	C C	T T
SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO	O. 46 O. 36	C C G T G T A T	G G	T T	G (G C	C C	A A	C C	C C	T T	C	G G	G G	A A	G G	A A	A A	A A	G G	T T	G G	G G	G G
SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO	O. 46 O. 36	T G C C C C	G G	T T	G (: A	T T	G G	A A	G G	C C	C	G G	C C	G G	C	G G	G G	C	C	T T	T T	T T
SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO SEQ. ID. NO	O. 46 O. 36	G A	G G	G G	G G	T C	T	G	G G	T T	G G	C C	G G	A A	G G	C C	C C	C C	T T	G G	C C	T T	G G	C C
SEQ. ID. N SEQ. ID. N SEQ. ID. N SEQ. ID. N	O. 46 O. 36	C T A G A G T T	A	A A	G G	C (C C	A A	G G	T T	G G	C C	C C	C C	G G	C C	G	T T	G G	G G	C C	T T	Ġ	T T

FIG. 11d.

SEQ SEQ SEQ	. ID. . ID.	NO.	. 46 . 36	C C	C	T T	G G	C T G	T T	C	A A	C C	C	C	G G	T T	T	C C	T T	G G	A A	G G	G G	A A	T T	G G	C C	С
SEQ. SEQ. SEQ.	ID. ID.	NO.	. 46 . 36	C C	G G	G G	G G	C A A	G G	C C	T T	G G	C C	T T	T T	G G	C C	T T	G G	C C	C C	A A	G G	C	C	A A	G G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	C C	C C	T T	TCCG	A A	A A	T T	G G	C C	C C	A A	G	C C	T T	T T	C C	A A	C C	C C	T T	G G	G G	G G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	G G	C C	C C	G A A C	G G	T T	G G	A A	T T	G G	G G	T T	T T	G G	G G	G G	G G	G G	G G	C C	C C	C C	T T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	A A	G G	A A	A G G C	T T	G G	T T	G G	G G	T T	G G	G G	C C	A A	G G	G G	C C	A A	G G	T T	G G	A A	G G	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	G G	G G	C C	T T C	G G	C C	T T	G G	A A	G G	G G	G G	T T	G G	C C	T T	A A	T T	C C	A A	C C	C	A A	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	G G	A A	G G	C C	T T	G G	G G	C C	C C	T T	C C	C C	T T	A A	C C	C C	C C	C C	A A	T T	C C	A A	G G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	A A	C C	T T	T T	T T	G G	C C	C C	T T G	C C G	C C	T T	A A	C C C	T T	T T	C C	C C	A A	G G	A A	G G	C C	C C

FIG. 11e.

SEQ SEQ SEQ	. ID . ID	. NO . NO	. 46 . 36	T	G	G	A A	C	C	C	T	7	G	G	A A	. A	C	A A	A A	C	A A	G G	C	C	G G	G G	A A	A A C
SEQ. SEQ. SEQ.	. ID. . ID.	. NO.	. 46 . 36	C	C	C	C	T T	G	G	T T	T	C	C	G G	T T	G G	A A	A A	T T	T T	C	T T	G	G G	G	A A	T G G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	G G	A A	G G	G G	T	T T	C	C C	G G	C C	A T T A	G G	C	A A	G G	C C	T T	T T	C C	C C	G G	G G	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 35	A A	G G	C C	G G	A A	G	А А	C	T T	G	C C	G G	G C C G	A A	G G	C C	C C	C C	A A	C C	T T	C C	T T	C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	C	G G	G	G G	C	T T	G	T T	G G	C C	C C	A C C	T T	T T	T T	G G	A A	G G	C C	A A	G G	G G	A A	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	C C	C	A A	A A	G G	A A	T T	C C	A A	T T	G G	C T T A	T T	T T	G G	T T	G G	G G	T T	C	A A	A A	T T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C	A A	G G	T T	G G	T T	A A	C C	G G	C	C C	A A	T T	G G	G G	C C	C	C C	A A	T T	G G	C C	G	C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C	C	A A	C C	A A	A A	C C	A A	T T	G	C	A A	C C	C	G G	T T	G	C	C	C	T	C	T T	G	C

FIG. 11f.

SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	C	C C	C C	A A	A A	C C	A A	C C	C C	A A	C C	C	C C	G G	G G	C C	T T	C C	T T	G G	T T	G G	A A	T C C	G G
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	C C	G G	A A	T T	G G	C C	G G	G G	C C	C	A A	G G	T T	T T	A A	A A	C	G G	G G	G G	C	G G	C C	C C T	G G
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	C C	C C	T T	C C	T T	A A	C C	A A	A A	G G	G G	A A	C C	T T	T T	T T	G G	T T	G G	C C	T T	C C	A A	C A A	C C
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	G G	T T	C C	A A	A A	G G	T T	T T	T T	G G	A A	T T	G G	C C	C C	C C	C C	C C	T T	T T	T T	C C	G G	A C C T	C C
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	C C	A A	G G	C C	T T	G G	A A	C C	A A	C C	C C	C C	A A	C C	A A	A A	T T	G G	A A	G G	G G	T T	C C	C C C G	G G
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	C C	T T	T T	T T	G G	A A	C C	C C	G G	C C	T T	T T	T T	G G	G G	T T	G G	A A	T T	G G	G G	T T	A A	A T T C	T T
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	G G	G	C C	C C	G	C C	T T	A A	C	A A	A A	C C	A A	T T	C C	T T	T T	C	A A	C C	C	T T	A A	T	C C
SEQ. I SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	T T	G G	C	G G	T T	G G	C C	A A	G G	G G	C C	A A	G G	T T	G G	G G	G G	C C	G G	C C	T T	A A	T T	C C	G G

FIG. 11g.

SEQ. SEQ. SEQ.	ID. ID.	NO.	46 36	C C	T T	A A	C C	C	A A	G G	A A	G A A C	G G	G G	T	G G	G G	G G	C C	T T	A A	C		G G	G G	O G	C	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	A A	A A	G G	G G	C C	T T	T T	A G G C	A A	C C	T T	C C	T T	G G	G G	A A	C C	A A	C C	C C	A A	G G	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	C C	A A	T T	C C	C C	C C	.А А	G T T G	G G	G G	G G	C C	C C	T T	C C	A A	C C	C C	C	T T	C C	A A	G G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	G G	G G	C C	C C	C C	C C	C C	T T T	G G	C C	C C	C C	G G	C C	C	T T	C	T T	C	C C	C C	T T	G G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	G G	T T	G G	A A	G G	C C	C	G C C C	T T	G G	C C	C C	T T	C C	C C	A A	G G	A A	A A	T T	G G	A A	G G	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	G G	A A	A A	G G	A A	G G	T T	C G A	T T	G G	C C	A A	G G	C C	C C	G G	G G	G G	C C	G G	A A	A A	G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C	T T	G G	C	T T	G	C	T T	G	G G	C C	T T	С	T T	G G	C C	A A	T T	T T	C C	C C	G G	T T	G G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C	A A	G G	C	C C	C	T T	A A	T T	G G	A A	G G	T T	A A	C C	C C	G G	A A	T T	T T	G	G G	A A	C C	C G

FIG. 11h.

SEQ. SEQ. SEQ.	ID. ID.	NO.	46 36	A A	A A	T T	T T	C C	A A	C	T T	T T	G	C C	G G	C	T T	G G	A A	T T	T T	G G	T T	G G	G	C	С	T
SEO. SEO. SEO. SEO.	ID. ID.	NO. NO.	46 36	G G	G G	.G G	C C	T T	A A	C C	T T	G G	G G	C C	C C	C C	A A	A A	T T	G G	C C	C C	A A	G G	C C	C	A T T C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	C C	T T	G G	G G	C C	T T	G G	C C	T T	T T	C C	G G	A A	A A	C C	T T	G G	c c	C C	C C	C	A A	G G G	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	G G	T T	A A	c c	A A	T T	C	C C	G G	C C	T T	G G	G G	G	G G	C C	G G	A A	T T	G G	C C	C C	A T T G	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	G G	C C	T T	G G	T T	G G	G G	G G	A A	C C	C C	T T	G G	T T	C C	A A	C C	C C	A A	T T	C C	G G	T C C A	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	G G	C C	C C	T T	C C	G G	G G	T T	G G	C C	C C	C	T T	G G	G G	C C	C C	A A	C C	C	C C	T T	G C C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	T T	G G	T T	G G	C C	T T	G G	G G	G G	T T	G G	T T	C C	T T	T T	T T	G G	T T	G G	C C	G G	G G	C C	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	A A	T T	G G	C C	C C	A A	C C	A A	C C	C C	A A	G G	T T	G G	G G	T T	C C	A A	A A	G G	G G	C C	C C

FIG. 111.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	C C	A A	G G	T G G C	T T	C	G G	G G	G G	A A	G G	C	T	C C	T	G G	C	T	A A	C	A A	T T	C	С С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	G G	C C	T T	C G G T	G G	G G	T T	G G	G G	T T	G G	T T	C C	T T	T T	C C	C C	T T	C	T T	G G	C C	T T	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	T T	G G	C C	C A A	T T	G G	A A	C C	C C	T T	T T	C C	A A	T T	C C	T T	T T	C C	A A	T T	T T	G G	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	A A	G G	C C	0 0 0	A A	T T	C C	C C	A A	C C	G G	G G	C C	A A	G G	T T	G G	T T	G G	T T	A A	C	C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	A A	C C	G G	0 0 0	C C	G G	T T	C C	T T	T T	G G	G G	T T	T T	T T	G G	G G	G G	C C	A A	C C	T T	G G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	T T	T T	C C	G T T A	C C	T T	G	T T	C C	T T	G	C C	T T	A A	C C	T T	C C	A A	G G	C	C C	C	T T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C	T T	C C	A A	C C	C	A A	A A	G G	A A	C C	C C	A A	A A	C C	C C	G G	C C	A A	T T	T T	G G	C	A A	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	C C	A A	T T	C C	T T	T T	C C	G G	G	T T	G G	G G	G	G G	C C	C	C	G G	G G	G G	A A	G G	G G	G G

FIG. 11j.

SEQ. SEQ. SEQ.	ID. ID.	NO.	46 36	T T	G G	C C	C	C C	A A	G G	C C	G G	G G	C C	C C	A A	C C	G G	C C	T T	T T	C	A A	T T	C C	A A	G G	Ŧ
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	C C	T T	G G	C C	C C	T T	C C	A A	C C	A A	G G	G G	T T	G G	G G	C C	C C	A A C	T T	C	T T	G G	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	G G	G G	C C	A A	C C	T T	T T	A A	T T	C C	T T	C C	G G	G G	G G	C C	C C	T A A T	G G	C C	T T	G	C C	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	T T	C C	G G	T T	G G	G G	T T	C C	G G	C C	C	T T	G G	G G	C C	T T	C G G	G G	T T	G G	G G	T T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	A A	G G	G G	C C	A A	C	C C	G G	G G	G G	C C	A A	C C	A A	G G	G G	C C	C A A G	A A	G G	G G	A A	G G	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	G G	C C	C C	C C	C C	C C	G G	A A	A A	C C	G G	G G	C C	G G	G G	G G	T A A G	G G	G G	T T	G G	G G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	A A	C C	A A	C C	T T	G G	C C	G G	C C	T T	G G	C C	А А	A A	C C	C	A A	C	C C	G G	C C	G G	A A	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	C C	A A	A A	G G	T T	A A	T T	G G	T T	T T	G G	G G	G G	C C	T T	C C	G G	C C	T T	G G	G G	C C	C C	T T

FIG. 11k.

SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	A A	C C	A A	A A	T T	G G	T T	G	C	T T	C	C C	T	C	A A	T T	C C	G	C C	G	C	T	C	T T T	G G
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	С	A A	C C	G G	C C	T T	T T	T T	A A	T T	G G	C C	C	T T	T T	C C	A A	A A	G G	A A	C	T T	C C	T G G C	C C
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	A A	A A	G G	T T	G G	C C	C C	C C	C	G G	A A	A A	A A	A A	C C	T T	T T	C C	A A	A A	C C	G G	A A	C G G G	G G
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	C C	C C	A A	A A	G G	T T	T T	C C	A A	T T	T T	G G	G G	C C	T T	T T	C C	A A	C	C	A A	T T	G	G T T A	A A
SEQ. I SEQ. I SEQ. I	D. D.	NO. NO.	46 36	C C	A A	C C	C C	A A	C C	C C	T T	G G	C C	A A	T T	C C	A A	T T	C C	T T	G G	G G	C C	T T	G G	G G	A C C G	A A
SEQ. I SEQ. I SEQ. I SEQ. I	D. D.	NO. NO.	46 36	T T	T T	C C	C C	T T	G G	C C	C	C C	A A	T T	C	T T	T T	C C	T T	A A	T T	G	T T	C C	A A	C C	T C C	T T
SEQ. I SEQ. I SEQ. I	D. D.	NO. NO.	46 36	C	C C	A A	G G	T T	G G	A A	C C	T T	A A	C C	C C	G G	G G	G G	T T	A A	C C	A A	G	A A	C C	C C	A A	C C
SEQ. I SEQ. I SEQ. I	ID. ID.	NO. NO.	46 36	C	A A	C C	C	A A	T T	G	T T	G G	C C	G G	T T	G G	T T	C C	A A	G G	T T	C C	A A	G G	C C	C C	T T	C C

FIG. 11L.

SEQ SEQ SEQ	. ID.	. NO . NO	. 46 . 36	A A	G	C	G G	T G G A	C	T T	C	C C	G G	T T	G	G	T T	G	C	T T	T T	G G	G G	C	T T	G G	C	C
SEQ. SEQ. SEQ.	. ID. . ID.	NO.	. 46 . 36	T T	C	T T	T T	G T T C	G G	C C	G G	C C	C C	C C	A A	A A	G G	C C	T T	G G	C C	A A	C C	A A	T T	C C	A A	T T
SEQ. SEQ. SEQ. SEQ.	. ID. . ID.	NO.	46 36	C C	C	T T	C C	A T T C	T T	C C	C C	A A	G G	C C	C C	G G	C	A A	G G	A A	A A	G G	A A	A A	C C	A A	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	T T	C C	G G	C A A T	G G	G G	A A	G G	G G	T T	G G	C C	G G	T T	T T	G G	C C	A A	G G	C C	A A	C C	C C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	G G	C C	G T T G	C C	A A	C	G G	C C	T T	T T	T T	C C	A A	A A	G G	G G	T T	G	G G	C C	T T	G G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	C C	G G	G G	C G A	C C	C C	A A	C C	G G	C C	T T	G G	C C	G G	C C	C C	G G	C C	A A	G G	C C	A A	A A	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	T T	C C	T T	С	C C	C C	G G	C C	A A	A A	G G	C C	G G	G G	T T	C C	C C	A A	G G	C C	A A	G G	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	T T	G	G	A A	G G	G G	C	T T	C	C	A A	C C	G G	G	G G	A A	T T	C	C	A A	C	C	C	C

FIG. I Im.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C	T T	C	C C	T T	C C	C C	T	C C	C C	C A A G	T T	C C	A A	G G	C C	A A	G G	C	A A	A A	G G	A A	G G	C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	A A	C C	A A	G G	C C	G G	A A	A A	G G	G A A G	C C	C C	C C	A A	T T	T T	C C	C C	C C	A A	С С	A A	G G	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	С	C C	G G	A A	G G	A A	G G	G G	C C	A A	0000	A A	A A	G G	C C	A A	G G	C	A A	G G	C C	A A	G	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	C C	T T	G G	G G	C C	C C	C C	T T	A A	A A G	C C	C C	C C	A A	G G	C	A A	A A	G G	A A	G G	C C	A A	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	G G	C C	A A	G G	C C	A A	G G	C	CCCG	C	C C	T T	G G	A A	C C	C C	C C	T T	C C	C	C	A A	C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	G G	C C	A A	G G	C C	A A	A A	C C	G G	G A A G	T T	C	T T	C C	A A	G G	C C	A A	G G	C	A A	G G	C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	46 36	C C	A A	G G	A A	T T	G G	C C	A A	A A	G G	C C	A A	G G	A A	A A	G G	G G	T	C	A A	T T	C	T T	T T	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G	G G	C	A A	G G	C C	G G	G G	C	A A	C	G G	G	T T	C	A A	C C	C C	T T	T T	C	T T	C C	A A	C C

FIG. 11n.

SEQ. SEQ. SEQ.	ID. ID.	NO NO	. 46 . 36	T T	G	A A	G	C C	T T	T T	T T	G G	A A	T T	G G	A A	G	C	C C	T T	C	A A	G G	A A	A A	G G	A A	G A A T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	. 46 . 36	C C	G G	C	C C C G	A A	T T	G G	G G	C C	C C	C C	A A	C C	G G	G G	G G	A A	A A	T T	T T	C C	T T	A A	C C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	. 46 . 36	C C	A A	C C	$G \cup G$	A A	G G	A A	A A	C C	T T	C C	C C	C C	T T	G G	G G	A A	G G	G G	C C	C	C C	A A	G G	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	A A	A A	T G G T	C C	A A	G G	C C	G G	A A	T T	A A	C C	G G	C C	T T	G G	A A	C C	C C	C C	G G	A A	C C	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	C C	A A	G	C C	C C	A A	T T	T T	A A	C C	T T	C	C C	C C	G G	C C	T T	G G	C C	A A	G G	T T	G G	C C
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	G G	G G	A G G A	A A	A A	A A	C C	G G	G G	A A	C C	T T	T T	A A	G G	A A	T T	C C	T T	G	A A	C C	C C	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	C C	C C	A A	G G	G G	A A	A A	A A	C C	A A	G G	G G	T T	C C	T T	G	C C	A A	A A	G G	G G	A A	C C	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	G	T T	G G	G G	G G	T T	G G	G G	A A	G G	A A	C C	C C	A A	G G	C C	G G.	G G	C	C	A A	G G	A A	G G

FIG. 110.

SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	T T	G	G G	A A	G G	G G	A A	C C	C C	C	T T	G G	A A	A A	G G	A A	G G	T T	T T	G	T T	C	$\circ \circ \circ \circ$	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C C	A A	G G	C C	A A	C C	T T	T T	G G	T T	A A	G G	T T	G G	T T	C C	C C	A A	G G	T T	T T	C	A A	G C C	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	A A	G G	C C	T T	T T	T T	G G	T T	C C	A A	T T	C C	A A	G G	T T	G G	G G	T T	G G	G G	A A	G C	T G G C	C C
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	G G	C C	A A	C C	T T	G G	T T	T T	A A	C C	A A	G G	A A	A A	A A	A A	C C	G G	T T	A A	G G	T T	A G G T	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A A	T T	T T	C C	A A	G A	C T	G G	G A	C	C T	G C	C T	C G	A G	T A	G G	A T	C C	T C	C A	T T	G C	A G A G	A T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	T G	C C	C G	A T	T G	C C	A T	T G	G C	G C	C T	G G	T A	G G	C	T G	G A	C G	C G	T A	G G	A G	G G C A	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G A	A A	G G	G G	A A	G A	G G	C C	C C	A C	A G	G G	G C	A G	A G	G A	C T	C C	C A	G A	G C	C G	G A		A G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T A	C G	A A	A T	C	G G	A A	C G	G C	A G	G G	A C	T A	C G	G C	A T	G C	C C	G G	G C	C A	A G	G G		T A

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SEQ. SEQ. SEQ. SEQ.	ID. ID. ID.	NO. NO. NO.	. 46 . 36 . 32	C C A	C A T	G A A	C G T	A C C	G G A	G G A	G G T	A A T	C A	A G T	A C C	G C T	C C G	G A	G C C	G C T	A G C	C G T	G G A C	C A C	0000	C A T	G T A	C A
SEQ. SEQ. SEQ.	ID.	NO.	36	Α	Α	G	С	T	G	С	Τ	G	С	T	G	С	T	С	G	G	G	Α	С	Α	G	G	C A C	G
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G A	G G	A A	C G	A T	G G	G G	A C	G A	A A	G G	A A	G G	T T	G A	G C	C G	A T	A T	G T	A A	G T	T C	A A A	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	T C	T A	T G	A A	T T	C G	A A	A G	G A	C A	A T	G C	A A	T T	G C	A C	G A	A T	A G	T G	C G	A T	G T C G	C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C G	A G	T A	G T	G A	G C	T T	C C	A T	G G	G A	A T	T G	A A	C A	T G	C A	T T	G A	A A	T A	G A	A G	G A G A	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A G	T C	A T	A T	A C	A A	G C	G C	G A	G A	C G	T C	T T	C G	A G	C T	C G	A T	A A	G T	C C	T A	G G	C G A A	T A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G C	T A	A T	T C	C T	A T	G C	A A	A C	C G	A G	T C	C C	T A	T T	C G	A C	C A	G G	G G	C	C C	A A	T T	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C A	A T	G C	G A	C G	C A	A G	T C	G C	A A	T T	C G	A G	G A	A C	G A	C C	C A	A C	T	G C	G A	A A	C G	A A

FIG. 11q.

SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	C A C T $T C C C$	CAAGA	A A G C A G A A T C C C A T A A G T A T G I T T G A A A	A C A A G T A G C A C A	ATGA
SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	G C A C G G C T	A A T A A	C A G G A T A A G G C T C A C A C A A T T T T C T A G T	T G C A C A A G T T C G	ATTA
SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	G T T C G T T G	G A G A A A T G T G	G G C T T C A G T T G A T G G A G A A G T C A A G T T	G T G G A G G T G T C T	A A G G G C T T
SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	T G T C T T G A	TGCTT	A C A T C T T T G A G A C A T A T G G A G A A C C	A T C C A T T A G A T G	ATGT
SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	A G A T A A A G	G C A A T A G T T T	T G A T C A G T A A A G A G T A T G G A A T C T C T T	T T T A T G T G A T C C	G A A T T G G A
SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	G A T C A T C C	C T G G A A G G A A	G A T C C C A A T C C A G T G C T A T C C C C T G G	G A A T G C G A T A G A	TATG
SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	A T A G G A G A	A C G A C A T A T C	G A G A A T C A A T T A T	A T C A A T C T G A C T	TATC
SEQ. ID. NO. 40 SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32	T G A C C A A A	T C T A C T A C T A	C	C T A T C T T G A C T T	TAAT

FIG. 11r.

SEQ. SEQ. SEQ.	. ID. . ID.	. NO. . NO.	46 36	G C	A G	G C C G	T	T T	G A	G G	A C	C T	C G	G A	C	G C	T	A T	G G	C	T C	G T	A A	C	C C	C T	T G	G
SEQ. SEQ. SEQ.	ID. ID.	NO.	46 36	C C	C T	G T A C	A C	C G	C C	T A	G A	C C	C A	T A	A G	C A	G T	C G	A T	A G	C C	A T	A T	G A	A G	T A	G G	T T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G T	C C	A T G A	T A	A G	G T	A C	G C	T C	T C	C A	G C	A C	G A	T C	C A	C G	C G	C G	A A	C T	C C	A A	C T	A C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	G A	T G A T	A T	T A	C C	A C	T C	C C	G T	A T	A T	T G	A A	C C	C T	C T	C A	T C	T A	T A	G A	A G	C T	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T T	A C	G C A C	A T	A T	A T	G T	T C	G A	T G	C A	A A	T T	T G	T G	T T	C C	A G	G A	A T	A G	T T	G A	G G	T G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C G	G G	A A G C	T C	G C	T A	A A	G A	G G	G G	G T	G C	C A	C G	A A	A G	A A	G G	G A	T A	C G	A A	G A	A A	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A T	G G	A G	A A	G T	A A	A C	A A	A C	T T	G G	G C	A T	T T	A T	C G	A A	C A	T A	G A	C T	T G	T T	T C	G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A C	A C	A T	A C	T T	G A	T T	C C	A A	C T	C G	T T	C T	T T	A C	T T	C A	A G	T T	G A	T G	T C	T G	C C	T T

FIG. 115.

SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	A T	G A	G T G T	A T	G G	C A	G A	C T	T A	T T	A G	G A	T T	G C	A A	A A	T	A T	T	C	A T	T C	C	A T	А
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G G	T A	A T G C	C T	T C	C A	G	T	G	G	A	G	T	С	Α	G	Α	C	Α	Α	T	G	Α	G	Α
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	Α	С	G C G	G	Α	Α	T	G	G	Α	G	G	Α	Α	Α	G	С	Α	Α	G	G	C	T	С	T
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	С	Т	G T T	Ţ	Α	G	Α	Α	С	Α	Α	T	Τ	Α	T	С	Α	C	Α	T	A.	С	С	C	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T	G	A G C	T	Т	С	C	Α	G	Α	Α	С	T	С	С	T	С	G	G	T	Ŧ	Α	T	T	С
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	T	G	G T A	T	C	T	T	Α	Α	Α	С	Α	Α	G	Α	Α	Α	G	Α	Т	C	T	Ţ	С	Ţ
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	Α	G	T A T	G	G	Α	G	A	Α	Α	Α	T	С	Α	Υ	G	Т	Α	Τ	T	С	С	C	Α	Ţ
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	С	T	Α	G	T	С	G	Α	С	T	Α	С	T	T	С	С	С	Α	G	Α	Α	Т	Α	T	G

FIG. 11t.

SUBSTITUTE SHEET (RULE 26)

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	Α	Ţ	G	G	Α	С	С	С	С	Α	G	Α	G	Α	G	Α	T	G	С	С	C	Α	G	C G G	C C A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36		G	C	С	С	G	Α	G	Α	А	T	T	С	Α	T	T	С	T	G	Α	Α	G	Α	Т	T G C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	C T C	Т	С	G	Т	G	G	Α	С	C	Т	G	Α	Α	C	C	C	Α	G	Α	C	Α	G	Τ	C G A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	Α	С	Α	Α	Α	Α	T	Т	Α	Т	С	T	Α	С	Т	С	C	C	Α	С	T	Ţ	С	C A A	C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G	T	G	С	G	Ç	С	Α	C	Α	G	Α	С	Α	С	С	G	Α	G	Α	Α	Т	Α	A T T	
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	С	G	С	Ţ	T	T	G	T	С	T	T	Ţ	G	C	Ţ	G	C	С	G	T	C	A	Α	G	0 0 0
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	Α	С	A	С	С	Α	T	С	С	Т	C	С	Α	G	T	T	G	Α	Α	С	C	T	G	Α	Α
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	46 36	G	G	Α	С	T	G	С	G	G	Т	C														

FIG. 1 IU.

SEQ. ID. NO. 40 GACAAAATTATCTACTCCCACTTCA SEQ. ID. NO. 46 CTCCTAGACACCCGCCCTGCCCTTC SEQ. ID. NO. 36 SEQ. ID. NO. 32 SEQ. ID. NO. 40 CGTGCGCCACAGACACCGAGAATAT SEQ. ID. NO. 46 CCTGGT SEQ. ID. NO. 36 SEQ. ID. NO. 32 SEQ. ID. NO. 40 CCGCTTTGTCTTTGCTGCCGTCAAG SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32 SEQ. ID. NO. 40 GACACCATCCTCCAGTTGAACCTGA SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32 SEQ. ID. NO. 40 AGGACTGCGGTCTGTTCTAA SEQ. ID. NO. 46 SEQ. ID. NO. 36 SEQ. ID. NO. 32

FIG. 1 /V.

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ClustalW Formatted Alignments

SEQ SEQ SEQ	. ID . ID.	. NO . NO	. 47 . 37	M M	G G	S S	L L	L	A A	L L	P P	Α	L L	L L	L L	L L	W W	G	A A	V V	A A	E	G G	P P	A A	K K	K K	L V V R
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO NO	. 47 . 37	L L	T T	L L	E E	G G	D D	L L	V V	Q L L L	G	G G	L L	F F	P P	v v	H H	Q Q	K K	G G	G G	P P	A A	E	D D	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	. 47 . 37	G G	P P	v v	N N	E E	H H	R R	G G	E I I E	QQ	R R	L L	E	A A	M M	L L	F F	A A	L L	D D	R R	I I	N N	R R	D D
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	47 37	P P	H H	L L	L L	P P	G G	V V	R R	L L	G G	A A	H H	I I	L L	D D	S S	C C	s s	K K	D D	T T	H H	A A	L L	E E
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	Q Q	A A	L L	D D	F F	v v	R R	A A	D S S E	L L	S S	R R	G G	A A	D D	G G	S S	R R	H H	I I	c C	P P	D D	G G	S S
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	Y Y	A A	T T	H H	G G	D D	A A	P P	A T T H	A A	I I	T T	G G	V V	1 1	G G	G G	S S	Y Y	S S	D D	v v	S	I I	Q Q
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	v v	A A	N N	L L	L L	R R	L L	F F	Q Q	I I	P P	Q Q	I I	S S	Y Y	A A	S S	T T	S S	A A	K K	L L	S S	D D	K K
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	S S	R R	Y Y	D D	Y Y	F F	A A	R R	T T	V V	P P	P P	D D	F F	F F	Q Q	A A	K K	A A	M M	A A	E	I I	L L	R R

FIG. 12a.

SEQ. SEQ. SEQ.	. <i>ID.</i> . ID.	NO.	. 47 . 37	F F	F F	N N	W W	T	Y Y	V V	s s	T	V V	A A	S S	E	G G	D D	Y Y	G	E	T	G	1	E	A A	F F	GEEK
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	. 47 . 37	L L	E E	A A	R R	A A	R R	N N	I I	C	V V	A A	T T	s s	E	K K	v v	G G	R R	A A	M M	S S	R R	A A	Q A A E	F F
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	E	G G	$\begin{array}{c} v \\ v \end{array}$	ν ν	R R	A A	L L	L L	Q Q	K K	P P	S S	A A	R R	v v	A A	V V	L L	F F	T T	R R	S S	Ε	N D D P	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	Ř R	E E	L L	L L	A A	A A	s s	Q Q	R R	L L	N N	A A	s s	F F	T T	w w	v v	A A	S S	D D	G G	w w	G G	Q A A W	L L
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	E	S S	V V	v v	A A	G G	s s	E	G G	A A	A A	E	G G	A A	I 1	T T	I I	E	L L	A A	S S	Y Y	P P	I I I A	\$ \$
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	D D	F F	A A	S S	Y Y	F F	Q Q	s s	L L	D D	P P	W W	N N	N N	s s	R R	N N	P P	w w	F F	R R	E	F F	T W W K	E E
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	Q Q	R R	F F	R R	C C	S S	F F	R R	Q Q	R R	D D	C C	A A	A A	H H	s s	L L	R R	A A	v V	P P	F F	E E	Q Q	E E
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	S S	K K	I I	M M	F F	v v	v v	N N	A A	V V	Y Y	A A	M M	A A	H H	A A	L L	H H	N N	M M	H H	R R	A A	L L	C C

FIG. 12b.

SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	P P	ソンスとの	T	T T	R R	L L	C C	D D	A A	M M	R R	P P	V V	N N	G G	R R	R R	L L	Y Y	K K	D D	F F	V V	L L	N
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	V V	Y K K I	F F	D D	A A	P P	F F	R R	P P	A A	D D	T T	H H	N N	E E	v v	R R	F F	D D	R R	F F	G G	D D	G G	I I
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	G G	N R R I	Y Y	7 N	I I	F F	T T	Y Y	L L	R R	A A	G G	s s	G G	R R	Y Y	R R	Y Y	Q Q	K K	V V	G G	Y Y	w w	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	E E	Q G G V	L L	T T	L L	D D	T T	S S	L L	I I	P P	w w	A A	S S	P P	s s	A A	G G	P P	L L	P P	A A	S S	R R	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	S S	M E E S	P P	C C	L L	Q Q	N N	E E	ν ν	K K	S S	v v	Q Q	P P	G G	E E	ν ν	C C	C C	W W	L L	C C	l I	P P	C C
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	Q Q	K P P K	Y Y	E	Y Y	R R	L L	D D	E E	F F	T T	C C	A A	D D	C C	G G	L L	G G	Y Y	W W	P P	7 7	A A	s s	L L
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	T T	G G	C C	F F	E E	L L	P P	Q Q	E E	Y Y	I I	R R	w w	G G	D D	A A	w w	A A	v v	G G	P P	V V	T T	I I	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	C	L L	G G	A A	L L	A A	T T	L L	F F	v v	L L	G G	v v	F F	V V	R R	H H	7 7	A A	T T	P P	V V	ν ν	K K	A A

FIG. 12c.

SEQ. SEQ. SEQ.	. ID. . ID.	NO.	. 47 . 37	S S	G	R R	E	L L	C	Y Y	I I	L L	L L	G G	G G	V V	F F	L L	C C	Y Y	C	M M	T T	F	I I	F F	I I	L A A L
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	K K	P P	S S	T T	A A	v v	C C	T T	L L	R R	R R	F L H	G G	L L	G G	T T	A A	F F	s s	V V	C C	Y Y	S S	A A	L L
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	L L	T T	K K	T T	Z Z	R R	I I	A A	R R	I I	F F	S G G Y	G G	A A	R R	E E	G G	A A	Q Q	R R	P P	R R	F F	1	S S
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	P P	A A	s s	Q Q	v v	A A	I I	C C	L L	A A	L L	P I I A	Տ Տ	G G	Q Q	L L	L L	I I	v v	v V	A A	W W	L L	v v	v v
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	E E	A A	P P	G G	T T	G G	K K	E	T T	A A	P P	V E E G	R R	R R	E E	V V	V V	T T	L L	R R	C C	7 7	H	R R	D D
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	A A	S S	M M	L L	G G	S S	L L	A A	Y Y	N N	V V	R L L	L L	I 1	A A	L L	C C	T T	L L	Y Y	A A	F F	K K	T T	R R
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	K K	C C	P P	E E	N N	F F	N N	E E	A A	K K	F F	I I	G G	F F	T T	M M	Y Y	T T	T T	C C	I I	I I	W W	L L	A A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	F F	L L	P P	I I	F F	Y Y	V V	T T	S S	S S	D D	Y Y	R R	V V	Q O	T T	T T	T T	M M	C	V V	S S	V V	\$ \$	L I.

FIG. 12d.

SUBSTITUTE SHEET (RULE 26)

SEQ. ID. NO. 4' SEQ. ID. NO. 4' SEQ. ID. NO. 3' SEQ. ID. NO. 3'	s G	G T A S V V S V V E A K	L G L G	C L	. F A	A P	K L K L	H H	I I I I	L L	F	Q Q	P P	Q Q	K K	N N	T T
SEQ. ID. NO. 4° SEQ. ID. NO. 4° SEQ. ID. NO. 3° SEQ. ID. NO. 3°	IE	V S L E V R E V R T S S	C S	T A T A	. A F	A i	F K	V .	A A A A	R R	A A	T T	L :	R R	R R	S S	N N
SEQ. 1D. NO. 41 SEQ. 1D. NO. 47 SEQ. 1D. NO. 33 SEQ. 1D. NO. 33	V S I	V R C R K R R K R C L F	S S S S	S L S L	. G C	i S i S	T G T G	S ·	T P	S S	\$ \$	S S	I I	S S	S S	K K	S S
SEQ. ID. NO. 4: SEQ. ID. NO. 4: SEQ. ID. NO. 3: SEQ. ID. NO. 3:	NS NS	K R S E D P E D P S R F	F P F P	Q P Q P	E R	Q	КQ КQ	Q	Q P Q P	L L	A A	L :	T (Q Q	Q Q	E E	Q Q
SEQ. ID. NO. 47 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	' Q Q (D P F Q P L Q P L V C N	T L T L	P Q P Q	Q Q) ·R) R	S Q S Q	Q (Q P Q P	R R	C C	K (Q Q	K K	V V	I I	F F
SEQ. ID. NO. 4: SEQ. ID. NO. 4: SEQ. ID. NO. 3: SEQ. ID. NO. 3:	7 G S (PLTGTV GTV LSE	T F T F	S L S L	S F	. D	E P E P	Q I	K N K N	A A	M M	A 1	H H	G G	N N	S S	T T
SEQ. ID. NO. 4: SEQ. ID. NO. 4: SEQ. ID. NO. 3: SEQ. ID. NO. 3:	HQ1	NSL NSL	E A E A	Q K Q K	S S S	D S	T L T L	T T	R H R H	Q Q	P P	L L	L L	P P	L L	Q Q	C
SEQ. ID. NO. 4 SEQ. ID. NO. 4: SEQ. ID. NO. 3: SEQ. ID. NO. 3:	GE GE	T D L T D L	D L D L	T V T V	'QE	E T	G L G L	Q Q	G P G P	ν ν	G	G	D D	Q Q	R R	P P	E

FIG. 12e.

SEQ. SEQ. SEQ.	ID. ID.	NO.	. 47 . 37	V V	E	D D	P P	E E	E E	L L	S S	P P	A A	L L	Λ. Λ.	V V	S	S S	S S	Q Q	S S	F F	\ \ \	i	S S	G	E G G K	G G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO.	47 37	S S	T T	V V	T T	E E	7 7	V V	V V	N N	S S	A M	A T	A L	M E	T S	L I	E M	S A	C	M C	A L	C S	CE	G L E I	S A
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	E K	E E	A A	K R	E R	A I	R N	R D	I E	N I	D E	E R	I Q	E L	R R	Q R	L D	R K	R R	D D	К А	R R	D R	A A E T	R L
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	R K	E L	L L	K L	L L	L G	L T	C L	G E	T S	G G	E K	S S	G T	K F	S I	T K	F Q	l M	K R	Q I	M T	R H	L I G V	1 5
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	H G	G Y	S S	C D	Y E	S D	D K	E R	D G	K F	R T	G K	F L	T V	K Y	L Q	V N	Y I	Q F	N T	1 A	F M	T O	S A A G	M M
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	Q I	A R	M A	I M	R D	A T	M L	D K	T I	L P	K Y	I K	P Y	Y E	K H	Y N	E K	H A	N H	K A	A O	H L	A V	M Q R L	L E
SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	V	R D	E V	V E	D K	V V	E	K A	V F	S E	A N	F P	E Y	N V	P D	Y A	V I	D K	A S	I L	K W	S N	L D	W P	N G
SEQ. SEQ. SEQ. SEQ.	ID. ID.	NO. NO.	47 37	D I	P Q	G E	I C	Q Y	E D	C R	Y R N	D R S	R E S	R Y V	R Q	E L L	Y S F	Q D L	L S	S T	D K	S Y	T Y	K !	Y N	Y	L I	N D

FIG. 12f.

SEQ. ID. NO. 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	D L D R V A	R V A D P A	D P A Y L P	Y L P T T Q Q D	T Q Q D O V L R	V L R V R V	L N D L D R V R V P T T P T T G I I Q A A R E F
SEQ. ID. NO. 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	G I I E Y P	EYP FDL	FDL QSV	Q S V I I F R M	I F R M 1 V D V	V D V G G Q	T T G I I E G G Q R S E R S E R R K F T C A T D
SEQ. ID. NO. 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	R R K W I H	WIH CFE	C F E N V T	NVTS SIMF	I M F E L V A	L V A L S E	S E R R K W L S E Y D Q Y D Q V L V L K D C G L
SEQ. ID. NO. 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	V L V	E S D	NEN	RMEE	ESKA	LFR	D Q V L V E T I I T Y P T Y P W F Q
SEQ. ID. NO, 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	WFQ	NSS	VIL	FLNK	KDL	LEE	Y P W F Q N K I M Y S H Y S H L V D
SEQ. ID. NO. 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	LVD	YFP	EYD	GPQR	DAQ	AAR	S H L V D Y E F 1 L K M L K M F V D
SEQ. ID. NO. 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	FVD	LNP	D S D	KIIY	SHF	$T \subset A$	TDTENI
SEQ. ID. NO. 41 SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33	R F V	FAA	V K D	SHF TILO	QLNL	KDC	N I R F V F G L F

FIG. 12g.

SEQ. ID. NO. 41 AAVKDTILQLNLKDCGLF SEQ. ID. NO. 47 SEQ. ID. NO. 37 SEQ. ID. NO. 33

FIG. 12h.

ClustalW Formatted Alignments

SEQ. SEQ.						G G																						
SEQ. SEQ.						C C																						
SEQ. SEQ.						G C																						
SEQ. SEQ.					C T	C A	T C	C T	A G	G C	A T	A G	G C	G T	T G	T C	G C	C G	C C	A T	G G	A C	T T	C G	A C	T T	A G	C C
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A C	C T	C C	C T	G G	C G	C	C G	T C	G C	G C	G	A G	A G	G G	G C	G C	G T	G G	C G	A G	T G	C C	A T	G G
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	G G	T	A C	C G	C	G G	G G	G G	G G	C	C G	T C	G C	A C	C C	T C	C	G G	G G	G C	A C	C G	C C	A C	G G
SEQ. SEQ.					T C	G C	A A	A G	G C	G A	C G	T C	A C	T C	C G	A C	A C	C G	T C	T T	C C	C T	T	G C	C A	C T	A C	G A
SEQ. SEQ.																												
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	C G	C G	G T	G G	G G	G C	G C	G A	A A	G G	C G	G G	C C	G A	A G	G C	G A	T T	G C	G G	T G	G G	G C	G	G C
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	C G	C G	C T	A G	A T	G G	G C	T	C	C	G C	C	A G	A C	G C	T	G T	C G	C G	T A	G A	G C	C T	C G	A G
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A C	C C	G A	G T	C	T G	C A	C G	T C	G A	G G	A A	C T	A C	G C	A G	T C	A A	T A	G C	G G	A A	C G	A T	C C
SEQ. SEQ.	ID. ID.	NO.	44 42	A A	C	C T	C	A C	G T	C G	С	G	С	С	C	T	G T 30	Α	С	C	G T	A C	A C	T	C C	T G	G A	C C

SUBSTITUTE SHEET (RULE 26)

SEQ. SEQ.	ID. ID.	NO. NO.	44 42	T C	C T	C G	A C	A G	G G	T C	C T	T C	T	A A	T	T G	T A	G C	A A	C	C G	C G	T A	G G	G T	A G	A C	A G
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A A	T C	G A	G A	G C	A G	A C	G A	G A	T A	T A	T G	T G	C G	C T	T	G G	A A	C A	G A	G G	G C	T C	G T	G T
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	G C	G T	A A	C C	C G	T A	C T	C G	C C	A A	G A	C T	T A	C A	T A	G A	G T	A A	C	G G	G G	A G	G C	C	C G
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	C A	G A	G C	G C	T A	G C	G T	A T	T G	T A	T T	C G	C G	G T	G G	T T	G T	T T	G G	A G	C A	C G	C G	C C	G G
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A T	C C	T	T G	C T	C C	A C	T A	C T	T C	G C	G G	T T	G C	G A	G C	C A	A T	G C	C C	T A	C T	C C	C A	G T
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	G T	A G	G C	C A	A G	T A	C G	T	G C	T C	A C	G T	T C	C	A A	G A	G G	G	C	C T	A G	G G	T A	G A	G T
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A C	G T	C G	A G	C T	C G	C	C A	C G	A C	A T	G T	C T	C	C T	C T	A T	C T	T G	G C	C T	C G	A C	G A	G A
SEQ. SEQ.																												
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SEQ.																									G		C	

FIG. 13b.

SEQ. ID. NO. 44 ATGAGCTCAAGCTCATCCACCACGA SEQ. ID. NO. 42 CTGAGGTGCGGAATGACCTGACTGG SEQ. ID. NO. 44 CAGCAAGTGTGATCCAGGCCAAGCC SEQ. ID. NO. 42 AGTTCTGTATGGCGAGGACATTGAG SEQ. ID. NO. 44 ACCAAGTACCTATATGAGCTGCTCT SEQ. ID. NO. 42 ATTTCAGACACCGAGAGCTTCTCCA SEQ. ID. NO. 44 A C A A C G A C C C T A T C A A G A T C A T C C T SEQ. ID. NO. 42 ACGATCCCTGTACCAGTGTCAAAAA SEQ. ID, NO. 44 TATGCCTGGCTGCAGCTCTGTCTCC SEQ. ID. NO. 42 GCTGAAGGGGAATGATGTGCGGATC SEQ. ID. NO. 44 ACGCTGGTGGCTGAGGCTGCTAGGA SEQ. ID. NO. 42 A T C C T T G G C C A G T T T G A C C A G A A T A SEQ. ID. NO. 44 TGTGGAACCTCATTGTGCTTTCCTA SEQ. ID. NO. 42 TGGCAGCAAAAGTGTTCTGTTGTGC SEQ. ID. NO. 44 TGGCTCCAGCTCACCAGCCCTGTCA SEQ. ID. NO. 42 ATACGAGGAGAACATGTATGGTAGT SEQ. ID. NO. 44 AACCGGCAGCGTTTCCCCACTTTCT SEQ. ID. NO. 42 AAATATCAGTGGATCATTCCGGGCT SEQ. ID. NO. 44 TCCGAACGCACCCATCAGCCACACT SEQ. ID. NO. 42 GGTACGAGCCTTCTTGGTGGGAGCA SEQ. ID. NO. 44 CCACAACCCTACCCGCGTGAAACTC SEQ. ID. NO. 42 GGTGCACACGGAAGCCAACTCATCC SEQ. ID. NO. 42 CGCTGCCTCCGGAAGAATCTGCTTG SEQ. ID. NO. 44 TTGCTACCATCCAGCAGACCACTGA SEQ. ID. NO. 42 CTGCCATGGAGGGCTACATTGGCGT

FIG. 13C. SUBSTITUTE SHEET (RULE 26)

SEQ. SEQ.														
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FIG. 13d. SUBSTITUTE SHEET (RULE 26)

SEQ. ID. NO. 44 GAGATTGTCATGCTGAATCCTGCCA SEQ. ID. NO. 42 ATTAAATTTACTCAATTTCAAGACA SEQ. ID. NO. 44 A T A C C C G C A G C A T T T C C A A C A T G A C SEQ. ID. NO. 42 GCAGGGAGGTGAAGGTGGGAGAGTA SEQ. ID. NO. 44 A T C C C A G G A A T T T G T G G A G A A C T A SEQ. ID. NO. 42 CAACGCTGTGGCCGACACACTGGAG SEQ. ID. NO. 44 ACCAAGCGACTGAAAAGACACCCTG SEQ. ID. NO. 42 A T C A T C A A T G A C A C C A T C A G G T T C C SEQ. ID. NO. 44 AGGAGACAGGAGGCTTCCAGGAGGC SEQ. ID. NO. 42 A A G G A T C C G A A C C A A A A G A C A A SEQ. ID. NO. 44 ACCGCTGGCCTATGATGCCATCTGG SEQ. ID. NO. 42 GACCATCATCCTGGAGCAGCTGCGG SEQ. ID. NO. 44 GCCTTGGCACTGGCCCTGAACAAGA SEQ. ID. NO. 42 AAGATCTCCCTACCTCTACAGCA SEQ. ID. NO. 44 CATCTGGAGGAGGCGGCCGTTCTGG SEQ. ID. NO. 42 TCCTCTCTGCCCTCACCATCCTCGG SEQ. ID. NO. 44 TGTGCGCCTGGAGGACTTCAACTAC SEQ. ID. NO. 42 GATGATCATGGCCAGTGCTTTTCTC SEQ. ID. NO. 44 AACAACCAGACCATTACCGACCAAA SEQ. ID. NO. 42 TTCTTCAACATCAAGAACCGGAATC SEQ. ID. NO. 44 TCTACCGGGCAATGAACTCTTCGTC SEQ. ID. NO. 42 AGAAGCTCATAAAGATGTCGAGTCC SEQ. ID. NO. 44 CTTTGAGGGTGTCTCTGGCCATGTG SEQ. ID. NO. 42 ATACATGAACAACCTTATCATCCTT SEQ. ID. NO. 44 G T G T T T G A T G C C A G C G G C T C T C G G A SEQ. ID. NO. 42 GGAGGGATGCTCTCCTATGCTTCCA

FIG /3e. SUBSTITUTE SHEET (RULE 26)

SEQ. ID. NO. 44 TGGCATGGACGCTTATCGAGCAGCT SEQ. ID. NO. 42 TATTTCTCTTTGGCCTTGATGGATC SEQ. ID. NO. 44 TCAGGGTGGCAGCTACAAGAAGATT SEQ. ID. NO. 42 C T T T G T C T C T G A A A G A C C T T T G A A SEQ. ID. NO. 44 GGCTACTATGACAGCACCAAGGATG SEQ. ID. NO. 42 A C A C T T T G C A C C G T C A G G A C C T G G A SEQ. ID. NO. 44 A T C T T T C C T G G T C C A A A A C A G A T A A SEQ. ID. NO. 42 TTCTCACCGTGGGCTACACGACCGC SEQ. ID. NO. 44 ATGGATTGGAGGGTCCCCCCAGCT SEQ. ID. NO. 42 TTTTGGGGCCATGTTTGCAAAGACC SEQ. ID. NO. 44 GACCAGACCCTGGTCATCAAGACAT SEQ. ID. NO. 42 TGGAGAGTCCACGCCATCTTCAAAA SEQ. ID. NO. 44 TCCGCTTCCTGTCACAGAAACTCTT SEQ. ID. NO. 42 ATGTGAAAATGAAGAAGATCAT SEQ. ID. NO. 44 TATCTCCGTCTCAGTTCTCCAGC SEQ. ID. NO. 42 CAAGGACCAGAAACTGCTTGTGATC SEQ. ID. NO. 44 CTGGGCATTGTCCTAGCTGTTGTCT SEQ. ID. NO. 42 GTGGGGGGCATGCTGCTGATCGACC SEQ. ID. NO. 44 GTCTGTCCTTTAACATCTACAACTC SEQ. ID. NO. 42 TGTGTATCCTGATCTGCTGGCAGGC SEQ. ID. NO. 44 ACATGTCCGTTATATCCAGAACTCA SEQ. ID. NO. 42 TGTGGACCCCCTGCGAAGGACAGTG SEQ. ID. NO. 44 CAGCCCAACCTGAACAACCTG SEQ. ID. NO. 42 GAGAAGTACAGCATGGAGCCGGACC SEQ. ID. NO. 44 CTGTGGGCTGCTCACTGGCTTTAGC SEQ. ID. NO. 42 CAGCAGGACGGGATATCTCCATCCG

FIG. 13f. SUBSTITUTE SHEET (RIJI F 26)

SEQ. ID. NO. 44 TGCTGTCTTCCCCCTGGGGCTCGAT SEQ. ID. NO. 42 CCCTCTCCTGGAGCACTGTGAGAAC SEQ. ID. NO. 44 GGTTACCACATTGGGAGGAACCAGT SEQ. ID. NO. 42 ACCCATATGACCATCTGGCTTGGCA SEQ. ID. NO. 44 TTCCTTTCGTCTGCCAGGCCCGCCT SEQ. ID. NO. 42 TCGTCTATGCCTACAAGGGACTTCT SEQ. ID. NO. 44 CTGGCTCCTGGGCCTGGGCTTTAGT SEQ. ID. NO. 42 CATGTTGTTCGGTTGTTTCTTAGCT SEQ. ID. NO. 44 CTGGGCTACGGTTCCATGTTCACCA SEQ. ID. NO. 42 TGGGAGACCCGCAACGTCAGCATCC SEQ. ID. NO. 44 AGATTTGGTGGGTCCACACGGTCTT SEQ. ID. NO. 42 CCGCACTCAACGACAGCAAGTACAT SEQ. ID. NO. 44 CACAAAGAAGGAAGAAAAGAAGAAG SEQ. ID. NO. 42 CGGGATGAGTGTCTACAACGTGGGG SEQ. ID. NO. 44 TGGAGGAAGACTCTGGAACCCTGGA SEQ. ID. NO. 42 ATCATGTGCATCATCGGGGCCGCTG SEQ. ID. NO. 44 AGCTGTATGCCACAGTGGGCCTGCT SEQ. ID. NO. 42 TOTCCTTCCTGACCCGGGACCAGCC SEQ. ID. NO. 44 GGTGGGCATGGATGTCCTCACTCTC SEQ. ID. NO. 42 CAATGTGCAGTTCTGCATCGTGGCT SEQ. ID. NO. 44 GCCATCTGGCAGATCGTGGACCCTC SEQ. ID. NO. 42 CTGGTCATCATCTTCTGCAGCACCA SEQ. ID. NO. 44 TGCACCGGACCATTGAGACATTTGC SEQ. ID. NO. 42 TCACCCTCTGCCTGGTATTCGTGCC SEQ. ID. NO. 44 CAAGGAGGAACCTAAGGAAAATATT SEQ. ID. NO. 42 GAAGCTCATCACCCTGAGAAAAC

F/G. 13g. SUBSTITUTE SHEET (RULE 26)

SEQ. SEQ.																												
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A G	G A	C T	A T	T C	T C	G A	C G	A T	G T	c c	T A	C C	C T	A C	G A	G G	A A	A A	G T	A C	T A	G G	A A	A A
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	T G	A A	C A	A A	T G	G A	G A	C G	T A	T T	G T	G C	C T	A A	T A	T A	T A	T C	C G	T	A C	T C	G A	G C	T C
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	T T	A C	C G	A G	A T	G C	G A	G C	G C	C A	T G	G T	C G	T	G G	C A	T A	G C	c c	T A	G A	C G	T C	G C	G A
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	G G	A C	A A	T C	C A	T T	T C	C C	C C	T G	T C	G C	C T	T G	T G	A A	T G	G G	A G	G C	A C	C T	C A	A C	A A
SEQ. SEQ.																												
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SEQ. SEQ.	ID. ID.	NO. NO.	44 42	C A	T	G T	T A	C A	C A	A C	G A	C G	C A	A A	G C	C C	A A	G C	G T	A A	T C	G C	C A	A A	G G	C A	C G	T C
SEQ. SEQ.																												
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FIG. 13h.
SUBSTITUTE SHFFT (BILLE 26)

SEQ. SEQ.																												
SEQ. SEQ.	. ID. . ID.	NO.	44	G C	T A	C C	G A	G A	A C	G A	G	C A	G G	C C	A C	G C	G T	A C	C T	A C	C G	C A	A A	T C	G A	A T	A G	G C
SEQ. SEQ.	ID.	NO. NO.	44 42	A A	C A	A A	G G	G A	G T	T C	C	A T	T A	C T	G A	A G	C A	C A	A G	A A	C T	A A	A T	C A	A A	A A	C C	G T
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A C	G T	G C	A C	G A	G G	A A	G A	A C	A A	G C	T A	C T	C C	C C	G A	G G	C C	T G	G T	T	T G	G G	G C	A T
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	G G	A T	A C	G C	G C	A T	G C	A C	A A	C G	C C	G T	T C	G C	A C	A C	C A	T T	G C	G C	A T	A C	A C	A A	G C
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A C	T A	C C	A G	T C	T C	G T	C A	T C	G C	A T	G C	A C	A C	A A	G T	A C	G C	G A	A T	G C	C G	G G	T A	G G
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	T G	C	T G	C T	T G	G G	A A	A C	C G	T C	G C	C A	G G	C C	C T	A G	T T	C G	A T	A C	C A	T G	C C	C	A C
SEQ. SEQ.																												
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	C C	G G	C C	C C	A A	C C	C A	C G	A A	C C	C A	G T	A G	C T	A G	C	C C	C A	C	C C	A C	G T	A C	A C	C T
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SEQ. SEQ.	ID. ID.	NO. NO.	44 42	T C	T C	T A	A A	T G	A G	Α	Α	G G //	С	С	C	G	G	G C	C G	C G	A A	T T	G C	A A	C A	T C	C G	T A
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SUBSTITUTE SHEET (RULE 26)

SEQ. ID. NO. 44 GGAGTCCATCATGGCGTGCTG SEQ. ID. NO. 42 CGAGATCGAGCGGCAGCTCCGCAGG SEQ. ID. NO. 44 A G C G A G G A G G C C A A G G A A G C C C G G C SEQ. ID. NO. 42 GACAAGCGGGACGCCGGCGGGAGC SEQ. ID. NO. 44 GGATCAACGAGAGATCGAGCGGCA SEQ. ID. NO. 42 TCAAGCTGCTGCTGCTCGGGACAGG SEQ. ID. NO. 44 GCTCCGCAGGGACAAGCGGGACGCC SEQ. ID. NO. 42 A G A G A G T G G C A A G A G T A C G T T T A T C SEQ. ID. NO. 44 CGCCGGGAGCTCAAGCTGCTGC SEQ. ID. NO. 42 AAGCAGATGAGAATCATCCATGGGT SEQ. ID. NO. 44 TCGGGACAGGAGAGTGGCAAGAG SEQ. ID. NO. 42 CAGGATACTCTGATGAAGATAAAG SEQ. ID. NO. 44 TACGTTTATCAAGCAGATGAGAATC SEQ. ID. NO. 42 GGGCTTCACCAAGCTGGTGTATCAG SEQ. ID. NO. 44 A T C C A T G G G T C A G G A T A C T C T G A T G SEQ. ID. NO. 42 AACATCTTCACGGCCATGCAGGCCA SEQ. ID. NO. 44 AAGATAAAAGGGGCTTCACCAAGCT SEQ. ID. NO. 42 TGATCAGAGCCATGGACACACTCAA SEQ. ID. NO. 44 GGTGTATCAGAACATCTTCACGGCC SEQ. ID. NO. 42 GATCCCATACAAGTATGAGCACAAT SEQ. ID. NO. 44 A T G C A G G C C A T G A T C A G A G C C A T G G SEQ. ID. NO. 42 AAGGCTCATGCACAATTAGTTCGAG SEQ. ID. NO. 44 A C A C A C T C A A G A T C C C A T A C A A G T A SEQ. ID. NO. 44 TGAGCACAATAAGGCTCATGCACAA SEQ. ID. NO. 42 TTTTGAGAATCCATATGTAGATGCA

FIG. 13j. SUBSTITUTE SHEET (R'JLE 26)

SEQ. ID. NO. 44 TTAGTTCGAGAAGTTGATGTGGAGA SEQ. ID. NO. 42 A T A A A G A G T T T A T G G A A T G A T C C T G SEQ. ID. NO. 44 AGGTGTCTGCTTTTGAGAATCCATA SEQ. ID. NO. 42 GAATCCAGGAATGCTATGATAGACG SEQ. ID. NO. 44 TGTAGATGCAATAAAGAGTTTATGG SEQ. ID. NO. 42 A C G A G A A T A T C A A T T A T C T G A C T C T SEQ. ID. NO. 44 AATGATCCTGGAATCCAGGAATGCT SEQ. ID. NO. 42 A C C A A A T A C T A T C T T A A T G A C T T G G SEQ. ID. NO. 44 A T G A T A G A C G A C G A G A A T A T C A A T T SEQ. ID. NO. 42 A C C G C G T A G C T G A C C C T G C C T A C C T SEQ. ID. NO. 44 A T C T G A C T C T A C C A A A T A C T A T C T T SEQ. ID. NO. 42 G C C T A C G C A A C A A G A T G T G C T T A G A SEQ. ID. NO. 44 AATGACTTGGACCGCGTAGCTGACC SEQ. ID. NO. 42 GTTCGAGTCCCCACAGGGATCA SEQ. ID. NO. 44 CTGCCTACCTGCCTACGCAACAAGA SEQ. ID. NO. 42 TCGAATACCCCTTTGACTTACAAAG SEQ. ID. NO. 44 TGTGCTTAGAGTTCGAGTCCCCACC SEQ. ID. NO. 42 TGTCATTTCAGAATGGTCGATGTA SEQ. ID. NO. 44 A C A G G G A T C A T C G A A T A C C C C T T T G SEQ. ID. NO. 42 GGGGGCCAAAGGTCAGAGAAAAAA SEQ. ID. NO. 44 ACTTACAAAGTGTCATTTTCAGAAT SEQ. ID. NO. 42 AATGGATACACTGCTTTGAAAATGT SEQ. ID. NO. 44 GGTCGATGTAGGGGGCCAAAGGTCA SEQ. ID. NO. 42 CACCTCTATCATGTTTCTAGTAGCG SEQ. ID. NO. 44 GAGAGAAGAAATGGATACACTGCT SEQ. ID. NO. 42 CTTAGTGAATATGATCAAGTTCTGG

FIG. 13 K. SUBSTITUTE SHEET (RULE 26)

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SEQ. SEQ.	ID. ID.	NO. NO.	44 42	A A	T C	G C	A T	T G	G A	G A	A C	C C	C C	C A	C G	A A	G C	A A	G G	A T	G G	A A	T C	G A	C A	C A	C A	A T
SEQ. SEQ.																												
SEQ. SEQ.																												
SEQ. SEQ.	ID. ID.	NO. NO.	44 42	G T	T C	G T	A T	C T	A G	A C	A T	G	T C	С	G	Т	С	Α	Α	C G	T G	C A	C C	C A	A C	C	T A	T T

FIG. 13 L. SUBSTITUTE SHEET (RULE 26)

SEQ. ID. NO. 44 CACGTGCGCCACAGACACCGAGAAT SEQ. ID. NO. 42 CCTCCAGTTGAACCTGAAGGGCTGC

SEQ. ID. NO. 44 A T C C G C T T T G T C T T T G C T G C C G T C A SEQ. ID. NO. 42 G G T C T G T A C

SEQ. ID. NO. 44 AGGACACCATCCTCCAGTTGAACCT SEQ. ID. NO. 42

SEQ. ID. NO. 44 GAAGGGCTGCGGTCTGTAC SEQ. ID. NO. 42

FIG. 13m.

ClustalW Formatted Alignments

SEQ. SEQ.	ID. ID.	NO. NO.	45 43	M M	L A	L S	L P	L R	L S	L S	A G	P Q	L P	F G	L P	R P	P P	P P	G P	A P	G P	G P	A P	Q A	T R	P L	N L	A L
SEQ. SEQ.	ID. ID.	NO. NO.	45 43	T L	S L	E	G L	C P	Q L	I L	I L	H P	P L	P A	W P	E G	G A	G W	I G	R W	Y A	R R	G G	L A	T P	R R	D P	Q P
SEQ. SEQ.	ID. ID.	NO. NO.	45 43	V P	K S	A S	I P	N P	F	L S	P I	V M	D G	Y L	E M	I P	E	Y T	v K	C E	R V	G A	E K	R G	E S	v I	V G	G R
SEQ. SEQ.	ID. ID.	NO. NO.	45 43	P G	K V	V L	R P	K A	C V	L E	A L	N A	G I	S E	w Q	T I	D R	M N	D E	T S	P L	S L	R R	C P	V Y	R F	I L	C D
SEQ. SEQ.												N D																
SEQ. SEQ.					V H	D L	F M	R V	C F	D G	P G	D V	F C	H P	L S	v v	G T	s s	S I	R I	S A	I E	C S	S L	Q Q	G G	Q W	W N
SEQ. SEQ.				S L																								
SEQ. SEQ.				F V	P P	M S	S D	G N	G A	w v	P N	G P	G A	Q I	A L	C K	Q L	P L	A K	V H	E Y	M Q	A W	L K	E R	D V	v G	N T
SEQ. SEQ.	ID. ID.	NO. NO.	45 43	S L	R T	R Q	D D	I V	L Q	P R	D F	Y S	E	L V	K R	L N	I D	H L	H T	D G	S V	K L	C Y	D G	P E	G D	Q I	A E
SEQ. SEQ.	ID.	NO. NO.	. 45 . 43	T I	K S	Y D	L T	Y E	E S	L F	L S	Y N	N D	D P	P C	I T	K S	I V	I K	L K	M L	P K	G G	C N	S D	s V	V R	S I
SEQ. SEQ.	. ID.	. NO: . NO:	. 45 . 43	T	L L	v G	A Q	E	A D	A Q	R N	M M	W A	N A	L K	I V	V F	C	s C	Y A	G Y	S E	S	Z Z	P M	A Y	L G	S S
SEQ. SEQ.	. ID . ID	. NO . NO	. 45 . 43	N K	R Y	Q	R W	F	P I	T P	F G	F W	R	T E	H P	P S	s w	A W	T	L Q	H V	N H	P T	T E	R A	V	K S	L S

FIG. 14a.

SUBSTITUTE SHEET (RULE 26)

SEQ. SEQ.																												
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SEQ. SEQ.																												
SEQ. SEQ.							A L																					
SEQ. SEQ.																												
SEQ. SEQ.																												
SEQ. SEQ.																												
SEQ. SEQ.																												
SEQ. SEQ.	ID. ID.	NO. NO.	45 43	s v	L G	G G	I M	V L	L L	A I	D	L	000	I	L	I	С	I W	Y Q	N A	s V	H D	V P	R L	Y R	I R	Q T	N V

FIG. 14b.

108/111

SEO. ID. NO. 45 SOPNLNNLTAVGCSLALAAVFPLGL SEQ. ID. NO. 43 EKYSMEPDPAGRDISIRPLLEHCEN SEQ. ID. NO. 45 DGYHIGRNQFPFVCQARLWLLGLGF SEQ. ID. NO. 43 THMTIWLGIVYAYKGLLMLFGCFLA SEQ. ID. NO. 45 SLGYGSMFTKIWWVHTVFTKKEEKK SEQ. ID. NO. 43 WETRNVSIPALNDSKYIGMSVYNVG SEQ. ID. NO. 45 EWRKTLEPWKLYATVGLLVGMDVLT SEO. ID. NO. 43 IMCIIGAAVS FLTRDQPNVQFCIVA SEQ. ID. NO. 45 LAIWQIVDPLHRTIETFAKEEPKED SEQ. ID. NO. 43 LVIIFCSTITLCLVFVPKLITLRTN SEQ. ID. NO. 45 I D V S I L P Q L E H C S S R K M N T W L G I F Y SEQ. ID. NO. 43 PDAATQNRRFQFTQNQKKEDSKTST SEQ.ID.NO.45 GYKGLLLLGIFLAYETKSVSTEKI SEQ. ID. NO. 43 SVTSVNQASTSRLEGLQSENHRLRM SEQ. ID. NO. 45 NDHRAVGMAIYNVAVLCLITAPVTM SEQ. ID. NO. 43 KITELDKDLEEVTMQLQDTPEKTTY *SEQ. ID. NO. 45* ILSSQQDAAFASLAIVFSSYITL SEQ. ID. NO. 43 IKQNHYQELNDILNLGNFTESTDGG SEQ.ID.NO.45 VVLFVPKMRRLITRGEWQSEAQDTM SEQ. ID. NO. 43 KAILKNHLDONPOLOWNTTEPSRTC SEQ.ID.NO.45 KTGSSTNNNEEEKSRLLEKENRELE SEQ. ID. NO. 43 KDPIEDINSPEHIQRRLSLQLPILH SEQ.ID.NO.45 KIIAEKEERVSELRHQLQSRQQLRS SEQ. ID. NO. 43 HAYLPSIGGVDASCVSPCVSPTASP SEQ. ID. NO. 45 RRHPPTPPEPSGGLPRGPPEPPDRL SEQ. ID. NO. 43 RHRHVPPS FRVMVSGLAAAMTLESI

FIG. 14c.

CHRCTTHITE CHEFT (RIH F 26)

SEQ. ID. NO. 45 S C D G S R V H L L Y K A A A M T L E S I M A C C SEQ. ID. NO. 43 MACCLSEEAKEARRINDEIEROLRR SEQ. ID. NO. 45 L S E E A K E A R R I N D E I E R Q L R R D K R D SEQ. ID. NO. 43 DKRDARRELKLLLGTGESGKSTFI SEQ. ID. NO. 45 ARRELKLLLLGTGESGKSTFIKOMR SEQ. ID. NO. 43 KOMRIIHGSGYSDEDKRGFTKLVYO SEQ. ID. NO. 45 IIHGSGYSDEDKRGFTKLVYQNIFT SEQ. ID. NO. 43 NIFTAMQAMIRAMDTLKIPYKYEHN SEQ.ID.NO.45 AMQAMIRAMDTLKIPYKYEHNKAHA SEQ. ID. NO. 43 KAHAQLVREVDVEKVSAFENPYVDA SEQ. ID. NO. 45 QLVREVDVEKVSAFENPYVDAIKSL SEQ. ID. NO. 43 IKSLWNDPGIQECYDRRREYOLSDS SEQ. ID. NO. 45 WNDPGIQECYDRRREYQLSDSTKYY SEQ. ID. NO. 43 TKYYLNDLDRVADPAYLPTQQDVLR SEQ. ID. NO. 45 LNDLDRVADPAYLPTQQDVLRVRVP SEQ. ID. NO. 43 VRVPTTGIIEYPFDLQSVIFRMVDV SEQ. ID. NO. 45 TTGIIEYPFDLQSVIFRMVDVGGQR SEQ. ID. NO. 43 GGQRSERRKWIHCFENVTSIMFLVA SEQ. ID. NO. 45 SERRKWIHCFENVTSIMFLVALSEY SEO. ID. NO. 43 L S E Y D Q V L V E S D N E N R M E E S K A L F R SEQ. ID. NO. 45 DQVLVESDNENRMEESKALFRTIIT SEQ. ID. NO. 43 TIITYPWFQNSSVILFLNKKDLLEE SEQ. ID. NO. 45 YPWFQNSSVILFLNKKDLLEEKIMY SEQ. ID. NO. 43 KIMYSHLVDYFPEYDGPQRDAQAAR SEQ. ID. NO. 45 SHLVDYFPEYDGPQRDAQAAREFIL SEQ. ID. NO. 43 EFILKMFVDLNPDSDKINYSHFTCA

FIG. 14d.

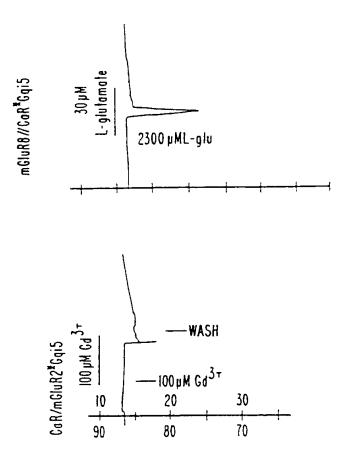
SUBSTITUTE SHEET (RULE 26)

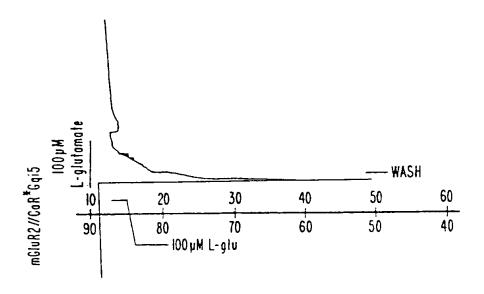
SEQ. ID. NO. 45 KM F V D L N P D S D K I I Y S H F T C A T D T E SEQ. ID. NO. 43 T D T E N I R F V F A A V K D T I L Q L N L K G C

SEQ. ID. NO. 45 NIR FV FAAVKD TILQLNLKGCGLY SEQ. ID. NO. 43 GLY

FIG. 14e.

FIG. 15.





SUBSTITUTE SHEET (RULE 26)

SEQUENCE LISTING

<110> NPS PHARMACEUTICALS, INC.

<120> G-PROTEIN FUSION RECEPTORS AND CHIMERIC GABA_R RECEPTORS

<130> 241/086-PCT

<140> TO BE ASSIGNED

<141> HEREWITH

<150> US 60/080,671

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<212> PRT

<213> Human

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Gln Asp Leu Lys Ser Arg Pro Glu Ser Val Glu Cys Ile Arg Tyr Asn 50 55 60

Phe Arg Gly Phe Arg Trp Leu Gln Ala Met Ile Phe Ala Ile Glu Glu 65 70 75 80

Ile Asn Ser Ser Pro Ala Leu Leu Pro Asn Leu Thr Leu Gly Tyr Arg 85 90 95

Ile Phe Asp Thr Cys Asn Thr Val Ser Lys Ala Leu Glu Ala Thr Leu 100 105 110

WO 99/51641

Ser	Phe	Val 115	Ala	Gln	Asn	Lys	11e 120	Asp	Ser	Leu	Asn	Leu 125	Asp	Glu	Ph
Cys	Asn 130	Cys	Ser	Glu	His	Ile 135	Pro	Ser	Thr	Ile	Ala 140	Val	Val	Gly	Al
Thr 145	Gly	Ser	Gly	Val	Ser 150	Thr	Ala	Val	Ala	Asn 155	Leu	Leu	Gly	Leu	Ph 16
Tyr	Ile	Pro	Gln	Val 165	Ser	Tyr	Ala	Ser	Ser 170	Ser	Arg	Leu	Leu	Ser 175	Ası
Lys	Asn	Gln	Phe 180	Lys	Ser	Phe	Leu	Arg 185	Thr	Ile	Pro	Asn	Asp 190	Glu	Hi
Gln	Ala	Thr 195	Ala	Met	Ala	Asp	Ile 200	Ile	Glu	Tyr	Phe	Arg 205	Trp	Asn	Trp
Val	Gly 210	Thr	Ile	Ala	Ala	Asp 215	Asp	Asp	Tyr	Gly	Arg 220	Pro	Gly	Ile	Glu
Lys 225	Phe	Arg	Glu	Glu	Ala 230	Glu	Glu	Arg	Asp	Ile 235	Cys	Ile	Asp	Phe	Se:
Glu	Leu	Ile	Ser	Gln 245	Tyr	Ser	Asp	Glu	Glu 250	Glu	Ile	Gln	His	Val 255	Val
Glu	Val	Ile	Gln 260	Asn	Ser	Thr	Ala	Lys 265	Val	Ile	Val	Val	Phe 270	Ser	Sei
Gly	Pro	Asp 275	Leu	Glu	Pro	Leu	Ile 280	Lys	Glu	Ile	Val	Arg 285	Arg	Asn	Ile
Thr	Gly 290	Lys	Ile	Trp	Leu	Ala 295	Ser	Glu	Ala	Trp	Ala 300	Ser	Ser	Ser	Let
Ile 305	Ala	Met	Pro	Gln	Tyr 310	Phe	His	Val	Val	Gly 315	Gly	Thr	Ile	Gly	Ph∈ 320
Ala	Leu	Lys	Ala	Gly 325	Gln	Ile	Pro	Gly	Phe 330	Arg	Glu	Phe	Leu	Lys 335	Lys
Val	His	Pro	Arg 340	Lys	Ser	Val	His	Asn 345	Gly	Phe	Ala	Lys	Glu 350	Phe	Trp
Glu	Glu	Thr 355	Phe	Asn	Cys	His	Leu 360	Gln	Glu	Gly	Ala	Lys 365	Gly	Pro	Leu
Pro	Val 370	Asp	Thr	Phe	Leu	Arg 375	Gly	His	Glu	Glu	Ser 380	Gly	Asp	Arg	Phe
Ser 385	Asn	Ser	Ser	Thr	Ala 390	Phe	Arg	Pro	Leu	Cys 395	Thr	Gly	Asp	Glu	Asr 400

3

Ile Ser Ser Val Glu Thr Pro Tyr Ile Asp Tyr Thr His Leu Arg Ile Ser Tyr Asn Val Tyr Leu Ala Val Tyr Ser Ile Ala His Ala Leu Gin 425 Asp Ile Tyr Thr Cys Leu Pro Gly Arg Gly Leu Phe Thr Asn Gly Ser Cys Ala Asp Ile Lys Lys Val Glu Ala Trp Gln Val Leu Lys His Leu 455 Arg His Leu Asn Phe Thr Asn Asn Met Gly Glu Gln Val Thr Phe Asp 470 475 Glu Cys Gly Asp Leu Val Gly Asn Tyr Ser Ile Ile Asn Trp His Leu Ser Pro Glu Asp Gly Ser Ile Val Phe Lys Glu Val Gly Tyr Tyr Asn 505 Val Tyr Ala Lys Lys Gly Glu Arg Leu Phe Ile Asn Glu Glu Lys Ile Leu Trp Ser Gly Phe Ser Arg Glu Val Pro Phe Ser Asn Cys Ser Arg 535 Asp Cys Leu Ala Gly Thr Arg Lys Gly Ile Ile Glu Gly Glu Pro Thr 545 550 555 Cys Cys Phe Glu Cys Val Glu Cys Pro Asp Gly Glu Tyr Ser Asp Glu Thr Asp Ala Ser Ala Cys Asn Lys Cys Pro Asp Asp Phe Trp Ser Asn 585

Thr Glu Pro Phe 610

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Glu Asn His Thr Ser Cys Ile Ala Lys Glu Ile Glu Phe Leu Ser Trp

600

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Ile	His	Pro 35	Pro	Trp	Glu	Gly	Gly 40	Ile	Arg	Tyr	Arg	Gly 45	Leu	Thr	Arg
Asp	Gln 50	Val	Lys	Ala	Ile	Asn 55	Phe	Leu	Pro	Vāl	Asp 60	Tyr	Glu	Ile	Glu
Tyr 65	Val	Cys	Arg	Gly	Glu 70	Arg	Glu	Val	Val	Gly 75	Pro	Lys	Val	Arg	Lys 80
Cys	Leu	Ala	Asn	Gly 85	Ser	Trp	Thr	Asp	Met 90	Asp	Thr	Pro	Ser	Arg 95	Cys
Val	Arg	Ile	Cys 100	Ser	Lys	Ser	Tyr	Leu 105	Thr	Leu	Glu	Asn	Gly 110	Lys	Val
Phe	Leu	Thr 115	Gly	Gly	Asp	Leu	Pro 120	Ala	Leu	Asp	Gly	Ala 125	Arg	Val	Asp
Phe	Arg 130	Cys	Asp	Pro	Asp	Phe 135	His	Leu	Val	Gly	Ser 140	Ser	Arg	Ser	Ile
Cys 145	Ser	Gln	Gly	Gln	Trp 150	Ser	Thr	Pro	Lys	Pro 155	His	Cys	Gln	Val	Asn 160
Arg	Thr	Pro	His	Ser 165	Glu	Arg	Arg	Ala	Val 170	Tyr	Ile	Gly	Ala	Leu 175	Phe
Pro	Met	Ser	Gly 180	Gly	Trp	Pro	Gly	Gly 185	Gln	Ala	Cys	Gln	Pro 190	Ala	Val
Glu	Met	Ala 195	Leu	Glu	Asp	Val	Asn 200	Ser	Arg	Arg	Asp	Ile 205	Leu	Pro	Asp
Tyr	Glu 210	Leu	Lys	Leu	Ile	His 215	His	Asp	Ser	Lys	Cys 220	Asp	Pro	Gly	Gln
Ala 225	Thr	Lys	Tyr	Leu	Tyr 230	Glu	Leu	Leu	Tyr	Asn 235	Asp	Pro	Ile	Lys	Ile 240
Ile	Leu	Met	Pro	Gly 245	Cys	Ser	Ser	Val	Ser 250	Thr	Leu	Val	Ala	Glu 255	Ala
Ala	Arg	Met	Trp 260	Asn	Leu	Ile	Val	Leu 265	Ser	Tyr	Gly	Ser	Ser 270	Ser	Pro
Ala	Leu	Ser 275	Asn	Arg	Gln	Arg	Phe 280	Pro	Thr	Phe	Phe	Arg 285	Thr	His	Pro
Ser	Ala 290	Thr	Leu	His	Asn	Pro 295		Arg	Val	Lys	Leu 300	Phe	Glu	Lys	Trp

Gly 305	Trp	Lys	Lys	Ile	Ala 310	Thr	Ile	Gln	Gln	Thr 315	Thr	Glu	Vai	Phe	Thr 320
Ser	Thr	Leu	Asp	Asp 325	Leu	Glu	Glъ	Arg	Val 330	Lys	Glu	Ala	Gly	11e 335	Glu
Ile	Thr	Phe	Arg 340	Gln	Ser	Phe	Phe	Ser 345	Asp	Pro	Ala	Val	Pro 350	Val	Lys
Asn	Leu	Lys 355	Arg	Gln	Asp	Ala	Arg 360	Ile	Ile	Val	Gly	Leu 365	Phe	Tyr	Glu
Thr	Glu 370	Ala	Arg	Lys	Val	Phe 375	Cys	Glu	Val	Tyr	Lys 380	Glu	Arg	Leu	Phe
Gly 385	Lys	Lys	Tyr	Val	Trp 390	Phe	Leu	Ile	Gly	Trp 395	Tyr	Ala	Asp	Asn	Trp
Phe	Lys	Ile	Tyr	Asp 405	Pro	Ser	Ile	Asn	Cys 410	Thr	Val	Asp	Glu	Met 415	Thr
Glu	Ala	Val	Glu 420	Gly	Hìs	Ile	Thr	Thr 425	Glu	Ile	Val	Met	Leu 430	Asn	Pro
Ala	Asn	Thr 435	Arg	Ser	Ile	Ser	Asn 440	Met	Thr	Ser	Gln	Glu 445	Phe	Val	Glu
Lys	Leu 450	Thr	Lys	Arg	Leu	Lys 455	Arg	His	Pro	Glu	Glu 460	Thr	Gly	Gly	Phe
Gln 465	Glu	Ala	Pro	Leu	Ala 470	Tyr	Asp	Ala	Ile	Trp 475	Ala	Leu	Ala	Leu	Ala 480
Leu	Asn	Lys	Thr	Ser 485	Gly	Gly	Gly	Gly	Arg 490	Ser	Gly	Val	Arg	Leu 495	Glu
Asp	Phe	Asn	Tyr 500	Asn	Asn	Gln	Thr	Ile 505	Thr	Asp	Gln	Ile	Tyr 510	Arg	Ala
Met	Asn	Ser 515	Ser	Ser	Phe	Glu	Gly 520	Val	Ser	Gly	His	Val 525	Val	Phe	Asp
Ala	Ser 530	Gly	Ser	Arg	Met	Ala 535	Trp	Thr	Leu	Ile	Glu 540	Gln	Leu	Gln	Gly
Gly 545	Ser	Tyr	Lys	Lys	Ile 550	Gly	Tyr	Tyr	Asp	Ser 555	Thr	Lys	Asp	Asp	Leu 560
Ser	Trp	Ser	Lys	Thr 565	Asp	Lys	Trp	Ile	Gly 570	Gly	Ser	Pro	Pro	Ala 575	Asp
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Pro	His	Leu 35	Pro	Arg	Pro	His	Ser 40	Arg	Val	Pro	Pro	His 45	Pro	Ser	Ser
Glu	Arg 50	Arg	Ala	Val	Tyr	Ile 55	Gly	Ala	Leu	Phe	Pro 60	Met	Ser	Gly	Gly
Trp 65	Pro	Gly	Gly	Gln	Ala 70	Cys	Gln	Pro	Ala	Val 75	Glu	Met	Ala	Leu	Glu 80
Asp	Val	Asn	Ser	Arg 85	Arg	Asp	Ile	Leu	Pro 90	Asp	Tyr	Glu	Leu	Lys 95	Leu
Ile	His	His	Asp 100	Ser	Lys	Cys	Asp	Pro 105	Gly	Gln	Ala	Thr	Lys 110	Tyr	Leu
Tyr	Glu	Leu 115	Leu	Tyr	Asn	Asp	Pro 120	Ile	Lys	Ile	Ile	Leu 125	Met	Pro	Gly
Cys	Ser 130	Ser	Val	Ser	Thr	Leu 135	Val	Ala	Glu	Ala	Ala 140	Arg	Met	Trp	Asn
Leu 145	Ile	Val	Leu	Ser	Tyr 150	Gly	Ser	Ser	Ser	Pro 155	Ala	Leu	Ser	Asn	Arg 160
Gln	Arg	Phe	Pro	Thr 165	Phe	Phe	Arg	Thr	His 170	Pro	Ser	Ala	Thr	Leu 175	His
Asn	Pro	Thr	Arg 180	Val	Lys	Leu	Phe	Glu 185	Lys	Trp	Gly	Trp	Lys 190	Lys	Ile
Ala	Thr	Ile 195	Gln	Gln	Thr	Thr	Glu 200	Val	Phe	Thr	Ser	Thr 205	Leu	Asp	Asp
Leu	Glu 210	Glu	Arg	Väl	Lys	Glu 215	Ala	Gly	Ile	Glu	Ile 220	Thr	Phe	Arg	Gln
Ser 225	Phe	Phe	Ser	Asp	Pro 230	Ala	Val	Pro	Val	Lys 235	Asn	Leu	Lys	Arg	Gln 240
Asp	Ala	Arg	Ile	Ile 245	Val	Gly	Leu	Phe	Tyr 250	Glu	Thr	Glu	Ala	Arg 255	Lys

Val Phe Cys Glu Val Tyr Lys Glu Arg Leu Phe Gly Lys Lys Tyr Val 265 Trp Phe Leu Ile Gly Trp Tyr Ala Asp Asn Trp Phe Lys Ile Tyr Asp Pro Ser Ile Asn Cys Thr Val Asp Glu Met Thr Glu Ala Val Glu Gly 295 His Ile Thr Thr Glu Ile Val Met Leu Asn Pro Ala Asn Thr Arg Ser Ile Ser Asn Met Thr Ser Gln Glu Phe Val Glu Lys Leu Thr Lys Arg 330 Leu Lys Arg His Pro Glu Glu Thr Gly Gly Phe Gln Glu Ala Pro Leu 340 345 Ala Tyr Asp Ala Ile Trp Ala Leu Ala Leu Ala Leu Asn Lys Thr Ser 360 Gly Gly Gly Gly Arg Ser Gly Val Arg Leu Glu Asp Phe Asn Tyr Asn 370 375 Asn Gln Thr Ile Thr Asp Gln Ile Tyr Arg Ala Met Asn Ser Ser Ser 395 390 Phe Glu Gly Val Ser Gly His Val Val Phe Asp Ala Ser Gly Ser Arg 410 405 Met Ala Trp Thr Leu Ile Glu Gln Leu Gln Gly Gly Ser Tyr Lys Lys 420 Ile Gly Tyr Tyr Asp Ser Thr Lys Asp Asp Leu Ser Trp Ser Lys Thr Asp Lys Trp Ile Gly Gly Ser Pro Pro Ala Asp Gln Thr Leu Val Ile 450 455 Lys Thr Phe Arg Phe Leu Ser Gln Lys

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- Pro Pro Pro Ser Ser Pro Pro Leu Ser Ile Met Gly Leu Met Pro Leu 50 55 60
- Thr Lys Glu Val Ala Lys Gly Ser Ile Gly Arg Gly Val Leu Pro Ala 65 70 75 80
- Val Glu Leu Ala Ile Glu Gln Ile Arg Asn Glu Ser Leu Leu Arg Pro
- Tyr Phe Leu Asp Leu Arg Leu Tyr Asp Thr Glu Cys Asp Asn Ala Lys
- Gly Leu Lys Ala Phe Tyr Asp Ala Ile Lys Tyr Gly Pro Asn His Leu 115 120 125
- Met Val Phe Gly Gly Val Cys Pro Ser Val Thr Ser Ile Ile Ala Glu 130 135 140
- Ser Leu Gln Gly Trp Asn Leu Val Gln Leu Ser Phe Ala Ala Thr Thr 145 150 155 160
- Pro Val Leu Ala Asp Lys Lys Lys Tyr Pro Tyr Phe Phe Arg Thr Val
- Pro Ser Asp Asn Ala Val Asn Pro Ala Ile Leu Lys Leu Leu Lys His 180 185 190
- Tyr Gln Trp Lys Arg Val Gly Thr Leu Thr Gln Asp Val Gln Arg Phe 195 200 205
- Ser Glu Val Arg Asn Asp Leu Thr Gly Val Leu Tyr Gly Glu Asp Ile 210 215 220
- Glu Ile Ser Asp Thr Glu Ser Phe Ser Asn Asp Pro Cys Thr Ser Val 225 230 235 240
- Lys Lys Leu Lys Gly Asn Asp Val Arg Ile Ile Leu Gly Gln Phe Asp 245 250 255
- Gln Asn Met Ala Ala Lys Val Phe Cys Cys Ala Tyr Glu Glu Asn Met 260 265 270
- Tyr Gly Ser Lys Tyr Gln Trp Ile Ile Pro Gly Trp Tyr Glu Pro Ser 275 280 285
- Trp Trp Glu Gln Val His Thr Glu Ala Asn Ser Ser Arg Cys Leu Arg 290 295 300
- Lys Asn Leu Leu Ala Ala Met Glu Gly Tyr Ile Gly Val Asp Phe Glu 305 310 315

Pro Leu Ser Ser Lys Gln Ile Lys Thr Ile Ser Gly Lys Thr Pro Gln 325 Gln Tyr Glu Arg Glu Tyr Asn Asn Lys Arg Ser Gly Val Gly Pro Ser 345 Lys Phe His Gly Tyr Ala Tyr Asp Gly Ile Trp Val Ile Ala Lys Thr Leu Gln Arg Ala Met Glu Thr Leu His Ala Ser Ser Arg His Gln Arg 375 Ile Gln Asp Phe Asn Tyr Thr Asp His Thr Leu Gly Arg Ile Ile Leu 390 385 Asn Ala Met Asn Glu Thr Asn Phe Phe Gly Val Thr Gly Gln Val Val Phe Arg Asn Gly Glu Arg Met Gly Thr Ile Lys Phe Thr Gln Phe Gln 420 Asp Ser Arg Glu Val Lys Val Gly Glu Tyr Asn Ala Val Ala Asp Thr Leu Glu Ile Ile Asn Asp Thr Ile Arg Phe Gln Gly Ser Glu Pro Pro

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<211> 583

<212> PRT

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Lys Asp Lys Thr Ile Ile Leu Glu Gln Leu Arg Lys Ile Ser Leu Pro

470

475

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Ser Gln Glu Tyr Ala His Ser Ile Arg Val Asp Gly Asp Ile Ile Leu

Gly Gly Leu Phe Pro Val His Ala Lys Gly Glu Arg Gly Val Pro Cys

Gly Glu Leu Lys Lys Glu Lys Gly Ile His Arg Leu Glu Ala Met Leu

Tyr	Ala	Ile	Asp	Gln 85	Ile	Asn	Lys	Asp	Pro 90	Asp	Leu	Leu	Ser	Asn 95	Ile
Thr	Leu	Gly	Val 100	Arg	Ile	Leu	Asp	Thr 105	Cys	Ser	Arg	Asp	Thr 110	Tyr	Ala
Leu	Glu	Gln 115	Ser	Leu	Thr	Phe	Val 120	Gln	Ala	Leu	Ile	Glu 125	Lys	Asp	Ala
Ser	Asp 130	Val	Lys	Cys	Ala	Asn 135	Gly	Asp	Pro	Pro	Ile 140	Phe	Thr	Lys	Pro
Asp 145	Lys	Ile	Ser	Gly	Val 150	Ile	Gly	Ala	Ala	Ala 155	Ser	Ser	Val	Ser	Ile 160
Met	Val	Ala	Asn	11e 165	Leu	Arg	Leu	Phe	Lys 170	Ile	Pro	Gln	Ile	Ser 175	Tyr
Ala	Ser	Thr	Ala 180	Pro	Glu	Leu	Ser	Asp 185	Asn	Thr	Arg	Tyr	Asp 190	Phe	Phe
Ser	Arg	Val 195	Val	Pro	Pro	Asp	Ser 200	Tyr	Gln	Ala	Gln	Ala 205	Met	Val	Asp
Ile	Val 210	Thr	Ala	Leu	Gly	Trp 215	Asn	Tyr	Val	Ser	Thr 220	Leu	Ala	Ser	Glu
Gly 225	Asn	Tyr	Gly	Glu	Ser 230	Gly	Val	Glu	Ala	Phe 235	Thr	Gln	Ile	Ser	Arg 240
Glu	Ile	Gly	Gly	Val 245	Cys	Ile	Ala	Gln	Ser 250	Gln	Lys	Ile	Pro	Arg 255	Glu
Pro	Arg	Pro	Gly 260	Glu	Phe	Glu	Lys	Ile 265	Ile	Lys	Arg	Leu	Leu 270	Glu	Thr
Pro	Asn	Ala 275	Arg	Ala	Val	Ile	Met 280	Phe	Ala	Asn	Glu	Asp 285	Asp	lle	Arg
Arg	Ile 290	Leu	Glu	Ala	Ala	Lys 295	Lys	Leu	Asn	Gln	Ser 300	Gly	His	Phe	Leu
Trp 305	Ile	Glγ	Ser	Asp	Ser 310	Trp	Gly	Ser	Lys	Ile 315	Ala	Pro	Val	Tyr	Gln 320
Gln	Glu	Glu	Ile	Ala 325	Glu	Gly	Ala	Vâl	Thr 330	Ile	Leu	Pro	Lys	Arg 335	Ala
Ser	Ile	Asp	Gly 340	Phe	Asp	Arg	Tyr	Phe 345	Arg	Ser	Arg	Thr	Leu 350	Ala	Asn
Asn	Arg	Arg	nzA	Vâl	Trp	Phe	Ala 360	Glu	Phe	Trp	Glu	Glu 365	Asn	Phe	Gly

11

Cys Lys Leu Gly Ser His Gly Lys Arg Asn Ser His Ile Lys Lys Cys 370 380

Thr Gly Leu Glu Arg Ile Ala Arg Asp Ser Ser Tyr Glu Gln Glu Gly 385 390 395 400

Lys Val Gln Phe Val Ile Asp Ala Val Tyr Ser Met Ala Tyr Ala Leu 405 410 415

His Asn Met His Lys Asp Leu Cys Pro Gly Tyr Ile Gly Leu Cys Pro 420 425 430

Arg Met Ser Thr Ile Asp Gly Lys Glu Leu Leu Gly Tyr Ile Arg Ala 435 440 445

Val Asn Phe Asn Gly Ser Ala Gly Thr Pro Val Thr Phe Asn Glu Asn 450 455 460

Gly Asp Ala Pro Gly Arg Tyr Asp Ile Phe Gln Tyr Gln Ile Thr Asn 465 470 475 480

Lys Ser Thr Glu Tyr Lys Val Ile Gly His Trp Thr Asn Gln Leu His 485 490 495

Leu Lys Val Glu Asp Met Gln Trp Ala His Arg Glu His Thr His Pro 500 505 510

Ala Ser Val Cys Ser Leu Pro Cys Lys Pro Gly Glu Arg Lys Thr 515 520 525

Val Lys Gly Val Pro Cys Cys Trp His Cys Glu Arg Cys Glu Gly Tyr 530 540

Asn Tyr Gln Val Asp Glu Leu Ser Cys Glu Leu Cys Pro Leu Asp Gln 545 550 555

Arg Pro Asn Met Asn Arg Thr Gly Cys Gln Leu Ile Pro Ile Ile Lys
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Leu Glu Trp His Ser Pro Trp 580

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Ala	Thr	Asn 35	Arg	Glu	Leu	Ser	Tyr 40	Leu	Leu	Leu	Phe	Ser 45	Leu	Leu	Cys
Cys	Phe 50	Ser	Ser	Ser	Leu	Phe 55	Phe	Ile	Gly	Glu	Pro 60	Gln	Asp	Trp	Thi
Cys 65	Arg	Leu	Arg	Gln	Pro 70	Ala	Phe	Gly	Ile	Ser 75	Phe	Val	Leu	Cys	11e
Ser	Cys	Ile	Leu	Val 85	Lys	Thr	Asn	Arg	Val 90	Leu	Leu	Val	Phe	Glu 95	Ala
Lys	Ile	Pro	Thr 100	Ser	Phe	His	Arg	Lys 105	Trp	Trp	Gly	Leu	Asn 110	Leu	Glr
Phe	Leu	Leu 115	Val	Phe	Leu	Cys	Thr 120	Phe	Met	Gln	Ile	Val 125	Ile	Cys	Va]
Ile	Trp 130	Leu	Tyr	Thr	Ala	Pro 135	Pro	Ser	Ser	Tyr	Arg 140	Asn	Gln	Glu	Leu
Glu 145	Asp	Glu	Ile	Ile	Phe 150	Ile	Thr	Cys	His	Glu 155	Gly	Ser	Leu	Met	Ala 160
Leu	Gly	Phe	Leu	lle 165	Gly	Tyr	Thr	Cys	Leu 170	Leu	Ala	Ala	Ile	Cys 175	Phe
Phe	Phe	Ala	Phe 180	Lys	Ser	Arg	Lys	Leu 185	Pro	Glu	Asn	Phe	Asn 190	Glu	Ala

Phe Ile Pro Ala Tyr Ala Ser Thr Tyr Gly Lys Phe Val Ser Ala Val

Lys Phe Ile Thr Phe Ser Met Leu Ile Phe Phe Ile Val Trp Ile Ser

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Glu Val Ile Ala Ile Leu Ala Ala Ser Phe Gly Leu Leu Ala Cys Ile 225 230 235 240

Phe Phe Asn Lys Ile Tyr Ile Ile Leu Phe 245 250

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Gln	Asn	Ser 35	Gln	Pro	Asn	Leu	Asn 40	Asn	Leu	Thr	Ala	Val 45	Gly	Cys	Ser
Leu	Ala 50	Leu	Ala	Ala	Val	Phe 55	Pro	Leu	Gly	Leu	Asp 60	Gly	Tyr	His	Ile
Gly 65	Arg	Asn	Gln	Phe	Pro 70	Phe	Val	Cys	Gln	Ala 75	Arg	Leu	Trp	Leu	Leu 80
Gly	Leu	Gly	Phe	Ser 85	Leu	Gly	Tyr	Gly	Ser 90	Met	Phe	Thr	Lys	Ile 95	Trp
Trp	Val	His	Thr 100	Val	Phe	Thr	Lys	Lys 105	Glu	Glu	Lys	Lys	Glu 110	Trp	Arg
Lys	Thr	Leu 115	Glu	Pro	Trp	Lys	Leu 120	Tyr	Ala	Thr	Val	Gly 125	Leu	Leu	Val
Gly	Met 130	Asp	Val	Leu	Thr	Leu 135	Ala	Ile	Trp	Gln	Ile 140	Val	Asp	Pro	Leu
His 145	Arg	Thr	Ile	Glu	Thr 150	Phe	Ala	Lys	Glu	Glu 155	Pro	Lys	Glu	Asp	Ile 160
Asp	Val	Ser	lle	Leu 165	Pro	Gln	Leu	Glu	His 170	Cys	Ser	Ser	Arg	Lys 175	Met
Asn	Thr	Trp	Leu 180	Gly	Ile	Phe	Tyr	Gly 185	Tyr	Lys	Gly	Leu	Leu 190	Leu	Leu
Leu	Gly	Ile 195	Phe	Leu	Ala	Tyr	Glu 200	Thr	Lys	Ser	Val	Ser 205	Thr	Glu	Lys
Ile	Asn 210	Asp	His	Arg	Ala	Val 215	Gly	Met	Ala	Ile	Tyr 220	Asn	Val	Ala	Val
Leu 225		Leu	Ile	Thr	Ala 230	Pro	Val	Thr	Met	Ile 235	Leu	Ser	Ser	Gln	Gln 240
Asp	Ala	Ala	Phe	Ala 245		Ala	Ser	Leu	Ala 250	Ile	Val	Phe	Ser	Ser 255	Tyr
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Gln Asn Ser Gln Pro Asn Leu Asn Asn Leu Thr Ala Val Gly Cys Ser 35 40 45

Leu Ala Leu Ala Ala Val Phe Pro Leu Gly Leu Asp Gly Tyr His Ile 50 55 60

Gly Arg Asn Gln Phe Pro Phe Val Cys Gln Ala Arg Leu Trp Leu Leu 65 70 75 80

Gly Leu Gly Phe Ser Leu Gly Tyr Gly Ser Met Phe Thr Lys Ile Trp 85 90 95

Trp Val His Thr Val Phe Thr Lys Lys Glu Glu Lys Lys Glu Trp Arg 100 105 110

Lys Thr Leu Glu Pro Trp Lys Leu Tyr Ala Thr Val Gly Leu Leu Val 115 120 125

Gly Met Asp Val Leu Thr Leu Ala Ile Trp Gln Ile Val Asp Pro Leu 130 135 140

His Arg Thr Ile Glu Thr Phe Ala Lys Glu Glu Pro Lys Glu Asp Ile 145 150 155 160

Asp Val Ser Ile Leu Pro Gln Leu Glu His Cys Ser Ser Arg Lys Met 165 170 175

Asn Thr Trp Leu Gly Ile Phe Tyr Gly Tyr Lys Gly Leu Leu Leu Leu 180 185 190

Leu Gly Ile Phe Leu Ala Tyr Glu Thr Lys Ser Val Ser Thr Glu Lys
195 200 205

Ile Asn Asp His Arg Ala Val Gly Met Ala Ile Tyr Asn Val Ala Val 210 215 220

Leu Cys Leu Ile Thr Ala Pro Val Thr Met Ile Leu Ser Ser Gln Gln 225 230 235 240

Asp Ala Ala Phe Ala Phe Ala Ser Leu Ala Ile Val Phe Ser Ser Tyr 245 250 255

Ile Thr Leu Val Val Leu Pne Val Pro Lys Met 260 265

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Ser Ala Phe Leu Phe Phe Asn Ile Lys Asn Arg Asn Gln Lys Leu Ile 20 25 30

Lys Met Ser Ser Pro Tyr Met Asn Asn Leu Ile Ile Leu Gly Gly Met 35 40 45

Leu Ser Tyr Ala Ser Ile Phe Leu Phe Gly Leu Asp Gly Ser Phe Val 50 55 60

Ser Glu Lys Thr Phe Glu Thr Leu Cys Thr Val Arg Thr Trp Ile Leu 65 70 75 80

Thr Val Gly Tyr Thr Thr Ala Phe Gly Ala Met Phe Ala Lys Thr Trp 85 90 95

Arg Val His Ala Ile Phe Lys Asn Val Lys Met Lys Lys Ile Ile 100 105 110

Lys Asp Gln Lys Leu Leu Val Ile Val Gly Gly Met Leu Leu Ile Asp 115 120 125

Leu Cys Ile Leu Ile Cys Trp Gln Ala Val Asp Pro Leu Arg Arg Thr 130 135 140

Val Glu Lys Tyr Ser Met Glu Pro Asp Pro Ala Gly Arg Asp Ile Ser 145 150 155 160

Ile Arg Pro Leu Leu Glu His Cys Glu Asn Thr His Met Thr Ile Trp 165 170 175

Leu Gly Ile Val Tyr Ala Tyr Lys Gly Leu Leu Met Leu Phe Gly Cys 180 185 190

Phe Leu Ala Trp Glu Thr Arg Asn Val Ser Ile Pro Ala Leu Asn Asp 195 200 205

Ser Lys Tyr Ile Gly Met Ser Val Tyr Asn Val Gly Ile Met Cys Ile 210 215 220

Ile Gly Ala Ala Val Ser Phe Leu Thr Arg Asp Gln Pro Asn Val Gln 225 230 235 240

Phe Cys Ile Val Ala Leu Val Ile Ile Phe Cys Ser Thr Ile Thr Leu 245 250 255

Cys Leu Val Phe Val Pro Lys Leu 260

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Phe Val Ile Val Thr Phe Val Arg Tyr Asn Asp Thr Pro Ile Val Arg 20 25 30

Ala Ser Gly Arg Glu Leu Ser Tyr Val Leu Leu Thr Gly Ile Phe Leu 35 40 45

Cys Tyr Ser Ile Thr Phe Leu Met Ile Ala Ala Pro Asp Thr Ile Ile 50 55 60

Cys Ser Phe Arg Arg Val Phe Leu Gly Leu Gly Met Cys Phe Ser Tyr 65 70 75 80

Ala Ala Leu Leu Thr Lys Thr Asn Arg Ile His Arg Ile Phe Glu Gln 85 90 95

Gly Lys Lys Ser Val Thr Ala Pro Lys Phe Ile Ser Pro Ala Ser Gln
100 105 110

Leu Val Ile Thr Phe Ser Leu Ile Ser Val Gln Leu Leu Gly Val Phe 115 120 125

Val Trp Phe Val Val Asp Pro Pro His Ile Ile Asp Tyr Gly Glu 130 135 140

Gln Arg Thr Leu Asp Pro Glu Lys Ala Arg Gly Val Leu Lys Cys Asp 145 150 155 160

Ile Ser Asp Leu Ser Leu Ile Cys Ser Leu Gly Tyr Ser Ile Leu Leu 165 170 175

PCT/US99/07333

Met Val Thr Cvs Thr Val Tyr Ala Ile Lys Thr Arg Gly Val Pro Glu 185

Thr Phe Asn Glu Ala Lys Pro Ile Gly Phe Thr Met Tyr Thr Thr Cys

Ile Ile Trp Leu Ala Phe Ile Pro Ile Phe Phe Gly Thr Ala Gln Ser 215

Ala Glu Lys Met Tyr Ile Gln Thr Thr Thr Leu Thr Val Ser Met Ser 230 235

Leu Ser Ala Ser Val Ser Leu Gly Met Leu Tyr Met Pro Lys Val Tyr 245 250

Ile Ile Ile Phe 260

WO 99/51641

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His Ala Phe Lys Val Ala Ala Arq Ala Thr Leu Arg Arq Ser Asn Val 20 25

Ser Arg Lys Arg Ser Ser Ser Leu Gly Gly Ser Thr Gly Ser Thr Pro

Ser Ser Ser Ile Ser Ser Lys Ser Asn Ser Glu Asp Pro Phe Pro Gln 55

Pro Glu Arg Gln Lys Gln Gln Gln Pro Leu Ala Leu Thr Gln Glu

Gln Gln Gln Gln Pro Leu Thr Leu Pro Gln Gln Gln Arg Ser Gln Gln

Gln Pro Arg Cys Lys Gln Lys Val Ile Phe Gly Ser Gly Thr Val Thr 100 105

Phe Ser Leu Ser Phe Asp Glu Pro Gln Lys Asn Ala Met Ala His Gly

Asn Ser Thr His Gln Asn Ser Leu Glu Ala Gln Lys Ser Ser Asp Thr 135 140

18

Leu Thr Arg His Gln Pro Leu Leu Pro Leu Gln Cys Gly Glu Thr Asp 145 150 155 160

Leu Asp Leu Thr Val Gln Glu Thr Gly Leu Gln Gly Pro Val Gly Gly 165 170 175

Asp Gln Arg Pro Glu Val Glu Asp Pro Glu Glu Leu Ser Pro Ala Leu 180 185 190

Val Val Ser Ser Ser Gln Ser Phe Val Ile Ser Gly Gly Ser Thr 195 200 205

Val Thr Glu Asn Val Val Asn Ser 210 215

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Met Lys Thr Gly Ser Ser Thr Asn Asn Glu Glu Glu Lys Ser Arg 20 25 30

Leu Leu Glu Lys Glu Asn Arg Glu Leu Glu Lys Ile Ile Ala Glu Lys
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Glu Glu Arg Val Ser Glu Leu Arg His Gln Leu Gln Ser Arg Gln Gln 50 55 60

Leu Arg Ser Arg Arg His Pro Pro Thr Pro Pro Glu Pro Ser Gly Gly 65 70 75 80

Leu Pro Arg Gly Pro Pro Glu Pro Pro Asp Arg Leu Ser Cys Asp Gly 85 90 95

Ser Arg Val His Leu Leu Tyr Lys 100

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Met Lys Thr Gly Ser Ser Thr Asn Asn Glu Glu Glu Lys Ser Arg
20 25 30

Leu Leu Glu Lys Glu Asn Arg Glu Leu Glu Lys Ile Ile Ala Glu Lys
35 40 45

Glu Glu Arg Val Ser Glu Leu Arg His Gln Leu Gln Ser Arg Gln Gln 50 55 60

Leu Arg Ser Arg Arg His Pro Pro Thr Pro Pro Glu Pro Ser Gly Gly 65 70 75 80

Leu Pro Arg Gly Pro Pro Glu Pro Pro Asp Arg Leu Ser Cys Asp Gly 85 90 95

Ser Arg Val His Leu Leu Tyr Lys

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Ile Thr Leu Arg Thr Asn Pro Asp Ala Ala Thr Gln Asn Arg Arg Phe 1 5 10 15

Gln Phe Thr Gln Asn Gln Lys Lys Glu Asp Ser Lys Thr Ser Thr Ser 20 25 30

Val Thr Ser Val Asn Gin Ala Ser Thr Ser Arg Leu Glu Gly Leu Gln 35 40 45

Ser Glu Asn His Arg Leu Arg Met Lys Ile Thr Glu Leu Asp Lys Asp 50 55 60

Leu Glu Glu Val Thr Met Gln Leu Gln Asp Thr Pro Glu Lys Thr Thr 65 70 75 80

Tyr Ile Lys Gln Asn His Tyr Gln Glu Leu Asn Asp Ile Leu Asn Leu 85 90 95

Gly Asn Phe Thr Glu Ser Thr Asp Gly Gly Lys Ala Ile Leu Lys Asn 100 105 110

20

His Leu Asp Gln Asn Pro Gln Leu Gln Trp Asn Thr Thr Glu Pro Ser 115 120 125

Arg Thr Cys Lys Asp Pro Ile Glu Asp Ile Asn Ser Pro Glu His Ile 130 135 140

Pro Ser Ile Gly Gly Val Asp Ala Ser Cys Val Ser Pro Cys Val Ser 165 170 175

Pro Thr Ala Ser Pro Arg His Arg His Val Pro Pro Ser Phe Arg Val 180 185 190

Met Val Ser Gly Leu 195

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Val Thr Ala Ala Thr Met Gln Ser Lys Leu Ile Gln Lys Gly Asn Asp 20 25 30

Arg Pro Asn Gly Glu Val Lys Ser Glu Leu Cys Glu Ser Leu Glu Thr 35 40 45

Asn Ser Lys Ser Ser Val Glu Phe Pro Met Val Lys Ser Gly Ser Thr 50 60

Ser 65

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Leu			Gln			_	Leu	Leu	Leu

Gly Pro Gly Glu Ser Gly Lys Ser Thr Phe Ile Lys Gln Met Arg Ile

Ile His Gly Val Gly Tyr Ser Glu Glu Asp Arg Arg Ala Phe Arg Leu 65 70 75 80

Leu Ile Tyr Gin Asn Ile Phe Val Ser Met Gln Ala Met Ile Asp Ala 85 90 95

Met Asp Arg Leu Gln Ile Pro Phe Ser Arg Pro Asp Ser Lys Gln His 100 \$105\$

Ala Ser Leu Val Met Thr Gln Asp Pro Tyr Lys Val Ser Thr Phe Glu 115 120 125

Lys Pro Tyr Ala Val Ala Met Gln Tyr Leu Trp Arg Asp Ala Gly Ile 130 135 140

Arg Ala Cys Tyr Glu Arg Arg Glu Phe His Leu Leu Asp Ser Ala 145 150 155 160

Val Tyr Tyr Leu Ser His Leu Glu Arg Ile Ser Glu Asp Ser Tyr Ile 165 170 175

Pro Thr Ala Gln Asp Val Leu Arg Ser Arg Met Pro Thr Thr Gly Ile

Asn Glu Tyr Cys Phe Ser Val Lys Lys Thr Lys Leu Arg Ile Val Asp 195 200 205

Val Gly Gly Gln Arg Ser Glu Arg Arg Lys Trp Ile His Cys Phe Glu 210 215 220

Asn Val Ile Ala Leu Ile Tyr Leu Ala Ser Leu Ser Glu Tyr Asp Gln 225 230 235 240

Cys Leu Glu Glu Asn Asp Gln Glu Asn Arg Met Glu Glu Ser Leu Ala 245 250 255

Leu Phe Ser Thr Ile Leu Glu Leu Pro Trp Phe Lys Ser Thr Ser Val 260 265 270

Ile Leu Phe Leu Asn Lys Thr Asp Ile Leu Glu Asp Lys Ile His Thr 275 280 285

Ser His Leu Ala Thr Tyr Phe Pro Ser Phe Glm Gly Pro Arg Arg Asp 290 295 300

22

Ala Glu Ala Ala Lys Ser Phe Ile Leu Asp Met Tyr Ala Arg Val Tyr 305 310 315 320

Ala Ser Cys Ala Glu Pro Gln Asp Gly Gly Arg Lys Gly Ser Arg Ala 325 330 335

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Arg Ser Val Phe Lys Asp Val Arg Asp Ser Val Leu Ala Arg Tyr Leu 355 360 365

Asp Glu Ile Asn Leu Leu 370

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Leu Glu Gln Lys Lys Gln Asp Arg Gly Glu Leu Lys Leu Leu Leu 25 40 45

Gly Pro Gly Glu Ser Gly Lys Ser Thr Phe Ile Lys Gln Met Arg Ile 50 60

Ile His Gly Ala Gly Tyr Ser Glu Glu Glu Arg Lys Gly Phe Arg Pro 65 70 75 80

Leu Val Tyr Gln Asn Ile Phe Val Ser Met Arg Ala Met Ile Glu Ala 85 90 95

Met Glu Arg Leu Gln Ile Pro Phe Ser Arg Pro Glu Ser Lys His His 100 105 110

Ala Ser Leu Val Met Ser Gln Asp Pro Tyr Lys Val Thr Thr Phe Glu 115 120 125

Lys Arg Tyr Ala Ala Ala Met Gln Trp Leu Trp Arg Asp Ala Gly Ile 130 135 140

Arg Ala Cys Tyr Glu Arg Arg Glu Phe His Leu Leu Asp Ser Ala 145 150 155 160

23

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Pro	Thr	Ala	Gln 180	Asp	Val	Leu	Arg	Ser 185	Arg	Met	Pro	Thr	Thr 190	Gly	Ιle
Asn	Glu	Tyr 195	Cys	Phe	Ser	Val	Gln 200	Lys	Thr	Asn	Leu	Arg 205	Ile	Vāl	Asp
Val	Gly 210	Gly	Gln	Lys	Ser	Glu 215	Arg	Lys	Lys	Trp	Ile 220	Eis	Cys	Phe	Glu
Asn 225	Val	Ile	Ala	Leu	Ile 230	Tyr	Leu	Ala	Ser	Leu 235	Ser	Glu	Tyr	Asp	Glr 240
Cys	Leu	Glu	Glu	Asn 245	Asn	Gln	Glu	Asn	Arg 250	Met	Lys	Glu	Ser	Leu 255	Ala
Leu	Pt₁e	Gly	Thr 260	Ile	Leu	Glu	Leu	Pro 265	Trp	Phe	Lys	Ser	Thr 270	Ser	Val
Ile	Leu	Phe 275	Leu	Asn	Lys	Thr	Asp 280	Ile	Leu	Glu	Glu	Lys 285	Ile	Pro	Thr
Ser	His 290	Leu	Ala	Thr	Tyr	Phe 295	Pro	Ser	Phe	Gln	Gly 300	Pro	Lys	Gln	Asp
Ala 305	Glu	Ala	Ala	Lys	Arg 310	Phe	Ile	Leu	Asp	Met 315	Tyr	Thr	Arg	Met	Tyr 320
Thr	Gly	Cys	Val	Asp 325	Gly	Pro	Glu	Gly	Ser 330	Lys	Lys	Gly	Ala	Arg 335	Ser
Arg	Arg	Leu	Phe 340	Ser	His	Tyr	Thr	Cys 345	Ala	Thr	Asp	Thr	Gln 350	Asn	Ile
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	agcagcagcc					
	agaaggtcat					2940 3000
	agaacgccat					3060
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				tggggcccaa		240
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DESCRIPTION AND DOCUMENTALLS

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WO 99/51641

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Pro	Ala	Ser	Gln	Val 725	Ala	Ile	Cys	Leu	Ala 730	Leu	Ile	Ser	Gly	Gln 735	Leu
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Ala 785	Leu	Cys	Thr	Leu	Tyr 790	Ala	Phe	Asn	Thr	Arg 795	Lys	Cys	Pro	Glu	Asn 600
Phe	Asn	Glu	Ala	Lys 805	Phe	Ile	Gly	Phe	Thr 810	Met	Tyr	Thr	Thr	Cys 815	lle

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- Arg Val Gln Thr Thr Thr Met Cys Val Ser Val Ser Leu Ser Gly Ser 835 840 845
- Val Val Leu Gly Cys Leu Phe Ala Pro Lys Leu His Ile Ile Leu Phe 850 855 860
- Gln Pro Gln Lys Asn Val Val Ser His Arg Ala Pro Thr Ser Arg Phe 865 870 875 880
- Gly Ser Ala Ala Arg Ala Ser Ser Ser Leu Gly Gln Gly Ser Gly 885 890 895
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53

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Val	Leu	Gly 35	Gly	Leu	Phe	Pro	Val 40	His	Gln	Lys	Gly	Gly 45	Pro	Ala	Glu
Asp	Cys 50	Gly	Pro	Val	Asn	Glu 55	His	Arg	Gly	Ile	Gln 60	Arg	Leu	Glu	Ala
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56

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Arg	Leu	Tyr	Lys 420	Asp	Phe	Val	Leu	Asn 425	Val	Lys	Phe	Asp	Ala 430	Pro	Phe
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WO 99/51641

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PCT/US99/07333

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Thr	Ala	Phe	Ser	Val 645	Cys	Tyr	Ser	Ala	Leu 650	Leu	Thr	Lys	Thr	Asn 655	Arg
Ile	Ala	Arg	Ile 660	Phe	Gly	Gly	Ala	Arg 665	Glu	Gly	Ala	Gln	Arg 670	Pro	Arg
Phe	Ile	Ser 675	Pro	Ala	Ser	Gln	Val 680		Ile	Cys	Leu	Ala 685	Leu	Ile	Ser
Gly	Gln 6 90		Leu	Ile	Val	Val 695		Trp	Leu	Val	Val 700	Glu	Ala	Pro	Gly
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Cys	Asn	His	Arg	Asp		Ser	Met	Leu	Gly 730	Ser	Leu	Ala	Tyr	Asn 735	Val

63

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- Glu Asp Lys Arg Gly Phe Thr Lys Leu Val Tyr Gln Asn Ile Phe Thr 1105 1110 1115 1120
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- Asn Arg Met Glu Glu Ser Lys Ala Leu Phe Arg Thr Ile Ile Thr Tyr 1285 1290 1295
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70

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- Gin Lys Val Ile Phe Gly Ser Gly Thr Val Thr Phe Ser Leu Ser Phe 945 950 955 960
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- Asn Ser Leu Glu Ala Gln Lys Ser Ser Asp Thr Leu Thr Arg His Gln 980 985 990
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1370

1350

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ε3

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91

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His Ala Leu Glu Gln Ala Leu Asp Phe Val Arg Ala Ser Leu Ser Arg

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Gly	Ala	Asp 115	Gly	Ser	Arg	His	Ile 120	Cys	Pro	Asp	Gly	Ser 125	Tyr	Ala	Th
His	Gly 130	Asp	Ala	Pro	Thr	Ala 135	Ile	Thr	Gly	Val	11e 140	Gly	Gly	Ser	Ту
Ser 145	Asp	Val	Ser	Ile	Gln 150	Val	Ala	Asn	Leu	Leu 155	Arg	Leu	Phe	Gln	11e
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Arg	Tyr	Asp	Tyr 180	Phe	Ala	Arg	Thr	Val 185	Pro	Pro	Asp	Phe	Phe 190	Gln	Ala
Lys	Ala	Met 195	Ala	Glu	Ile	Leu	Arg 200	Phe	Phe	Asn	Trp	Thr 205	Tyr	Val	Sei
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- Phe Val Phe Ala Ala Val Lys Asp Thr Ile Leu Gln Leu Asn Leu Lys 1380 1385 1390

Asp Cys Gly Leu Phe 1395

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CLASSIFICATION OF SUBJECT MATTER CO7K14/705 012N15/12 012N15/62 A61K38/17 According to International Patent Classification (IPC) or to both national classification and (PC B. FIELDS SEARCHED $\textbf{Minimum documentation searched} \ \ \text{classification system to lowed by classification symbols};$ Occumentation, searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages 13-28, WO 97 05252 A (NPS PHARMA INC) 33,38,40 13 February 1997 (1997-02-13) cited in the application see the whole document and specially pages 59-60. WO 97 46675 A (NOVARTIS AG) Υ 13-28, 11 December 1997 (1997-12-11) 33,38,39 cited in the application page 2, last paragraph - page 3, paragraph 1 page 50-56 page 62-67 page 86-90 -/--Patent family members are listed in annex Further documents are listed in the continuation of box C χ Special categories of cited documents "T" later document published after the international filing date or priority date and not in conflict with the application but "A" document defining the general state of the lart which is not cited to understand the principle or theory underlying the considered to be of particular relevance invention "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention frling date cannot be considered novel or cannot be considered. to "E" document which may throw doubts on pnority claim(s) or which is cited to establish the publication date of another involve an inventive step when the document is taken alone "Y" document of particular relevance: the claimed invention citation or other special reason (as specified) cannot be considered to involve an inventive step when the document is combined with one or more other, such docu-"O" document reterring to an oral disclosure, use, exhibition or other means ments, such combination being obvious to all person skilled "P" gocument published prior to the international filing date but "&" document member of the same patent family later than the phority date claimed Date of mailing of the international search report Date of the actual completion of the international search 13 September 1999 28/09/1999 Name and making address of the ISA Authorized officer European Patent Office P.E. 5818 Patentiaan 2 NL - 2280 HV Alswit Tei (+31-70) 340-2040, Tx 31 651 epc ni. Mateo Rosell, A.M. Fax: (+31-70) 340-3016

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		KENNETH A (US): LAZ THOMAS M (US); SYNAPTIC P) 29 April 1999 (1999-04-29) abstract page 105-109; table 3 figures 2A-D	13-28, 38.39
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